

10/513699

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NEWS 5 APR 28 IMSRESEARCH reloaded with enhancements
NEWS 6 MAY 30 INPAFAMDB now available on STN for patent family searching
NEWS 7 MAY 30 DGENE, PCTGEN, and USGENE enhanced with new homology sequence search option
NEWS 8 JUN 06 EPFULL enhanced with 260,000 English abstracts
NEWS 9 JUN 06 KOREPAT updated with 41,000 documents
NEWS 10 JUN 13 USPATFULL and USPAT2 updated with 11-character patent numbers for U.S. applications
NEWS 11 JUN 19 CAS REGISTRY includes selected substances from web-based collections
NEWS 12 JUN 25 CA/Caplus and USPAT databases updated with IPC reclassification data
NEWS 13 JUN 30 AEROSPACE enhanced with more than 1 million U.S. patent records
NEWS 14 JUN 30 EMBASE, EMBAL, and LEMBASE updated with additional options to display authors and affiliated organizations
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NEWS 16 JUN 30 STN AnaVist enhanced with database content from EPFULL
NEWS 17 JUL 28 CA/Caplus patent coverage enhanced
NEWS 18 JUL 28 EPFULL enhanced with additional legal status information from the epoline Register
NEWS 19 JUL 28 IFICDB, IFIPAT, and IFIUDB reloaded with enhancements
NEWS 20 JUL 28 STN Viewer performance improved
NEWS 21 AUG 01 INPADOCDB and INPAFAMDB coverage enhanced
NEWS 22 AUG 13 CA/Caplus enhanced with printed Chemical Abstracts page images from 1967-1998
NEWS 23 AUG 15 CAOLD to be discontinued on December 31, 2008
NEWS 24 AUG 15 CAplus currency for Korean patents enhanced
NEWS 25 AUG 25 CA/Caplus, CASREACT, and IFI and USPAT databases enhanced for more flexible patent number searching
NEWS 26 AUG 27 CAS definition of basic patents expanded to ensure comprehensive access to substance and sequence information

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NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3,
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	ENTRY	SESSION
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STRUCTURE FILE UPDATES: 14 SEP 2008 HIGHEST RN 1049627-95-3
DICTIONARY FILE UPDATES: 14 SEP 2008 HIGHEST RN 1049627-95-3

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TSCA INFORMATION NOW CURRENT THROUGH July 5, 2008.

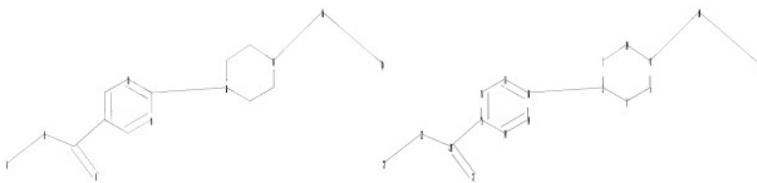
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REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stnqgen/stndoc/properties.html>

=>
Uploading C:\Program Files\Stnexp\Queries\10506998allow.str

10/513699



chain nodes :

10 11 20 21 22 23

ring nodes :

1 2 3 4 5 14 15 16 17 18 19 24

chain bonds :

2-18 4-10 10-11 15-20 20-21 20-22 22-23

ring bonds :

1-2 1-5 2-3 3-24 4-5 4-24 14-15 14-19 15-16 16-17 17-18 18-19

exact/norm bonds :

1-2 1-5 2-3 2-18 3-24 4-10 4-5 4-24 10-11 20-21 20-22 22-23

exact bonds :

15-20

normalized bonds :

14-15 14-19 15-16 16-17 17-18 18-19

isolated ring systems :

containing 1 :

G1:C,N

G2:Ak,NH2,NO2

G3:O

G4

G5:C,N,Zn,H

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 10:CLASS 11:Atom 14:Atom 15:Atom

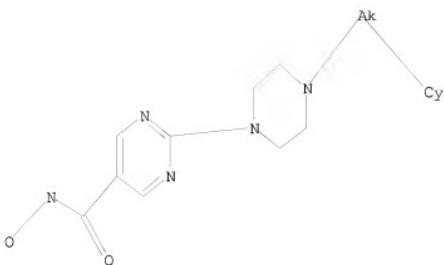
16:Atom 17:Atom 18:Atom 19:Atom 20:CLASS 21:CLASS 22:CLASS 23:CLASS 24:Atom

L1 STRUCTURE UPLOADED

=> d 11

L1 HAS NO ANSWERS

L1 STR



G1 C,N
 G2 Ak,NH₂,NO₂
 G3 O
 G4
 G5 C,N,Zn,H

Structure attributes must be viewed using STN Express query preparation.

=> s l1 full
 FULL SEARCH INITIATED 16:01:46 FILE 'REGISTRY'
 FULL SCREEN SEARCH COMPLETED - 1679 TO ITERATE

100.0% PROCESSED 1679 ITERATIONS 113 ANSWERS
 SEARCH TIME: 00.00.01

L2 113 SEA SSS FUL L1

=> file caplus			
COST IN U.S. DOLLARS	SINCE FILE	TOTAL	
FULL ESTIMATED COST	ENTRY	SESSION	
	178.36	178.57	

FILE 'CAPLUS' ENTERED AT 16:01:51 ON 15 SEP 2008
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FILE COVERS 1907 - 15 Sep 2008 VOL 149 ISS 12
FILE LAST UPDATED: 14 Sep 2008 (20080914/ED)

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reclassification data for the second quarter of 2008.

Effective October 17, 2005, revised CAS Information Use Policies apply.
They are available for your review at:

<http://www.cas.org/legal/infopolicy.html>

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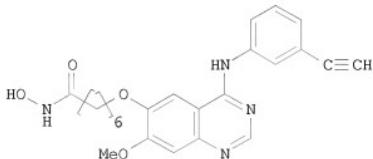
L3 11 L2

=> d ibib abs hitstr tot

L3 ANSWER 1 OF 11 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2008:353001 CAPLUS
 DOCUMENT NUMBER: 148:355828
 TITLE: Multi-functional small molecules as anti-proliferative agents and their preparation
 INVENTOR(S): Cai, Xiong; Qian, Changgeng; Gould, Stephen; Zhai, Haixiao
 PATENT ASSIGNEE(S): Curis, Inc., USA
 SOURCE: PCT Int. Appl., 494pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008033747	A2	20080320	WO 2007-US77971	20070910
WO 2008033747	A9	20080724		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA				
US 20080221132	A1	20080911	US 2007-852458	20070910
PRIORITY APPLN. INFO.:			US 2006-843590P	P 20060911
			US 2007-895889P	P 20070320

OTHER SOURCE(S): MARPAT 148:355828
 GI



A—B—C I

II

AB The invention relates to the compns., methods, and applications of an approach to selective inhibition of several cellular or mol. targets with a single small mol. More specifically, the present invention relates to multi-functional small mols. of formula I wherein one functionality is capable of inhibiting histone deacetylases (HDAC) and the other functionality is capable of inhibiting a different cellular or mol. pathway involved in aberrant cell proliferation, differentiation or

survival. Compds. of formula I wherein A is a pharmacophore of an anticancer agent capable of inhibiting at least one cellular or mol. pathway involved in the aberrant cell proliferation, differentiation or survival; B is a linker; C is a zinc-binding moiety; and their geometrical isomers, enantiomers, diastereoisomers, racemates, pharmaceutically acceptable salts, prodrugs and solvates thereof, are claimed. Example compound II was prepared by a multistep procedure (procedure given). All the invention compds. were evaluated for their antiproliferative activity (some data given).

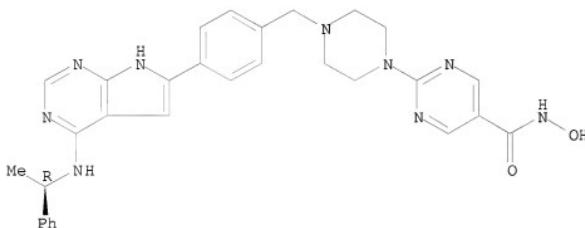
IT 1011716-90-7P

RL: PAC (Pharmacological activity); PKT (Pharmacokinetics); PRPH (Prophetic); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (prophetic starting material; preparation of multi-functional small mols. as antiproliferative agents)

RN 1011716-90-7 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[4-[4-[(1R)-1-phenylethyl]amino]-7H-pyrrolo[2,3-d]pyrimidin-6-yl]phenyl]methyl]-1-piperazinyl- (CA INDEX NAME)

Absolute stereochemistry.



L3 ANSWER 2 OF 11 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2008:351928 CAPLUS
 DOCUMENT NUMBER: 148:355814
 TITLE: Preparation of (aralkylamino)(phenyl)pyrrolo[2,3-d]pyrimidine derivatives for use as protein tyrosine kinase (PTK) inhibitors
 INVENTOR(S): Cai, Xiong; Qian, Changgeng; Gould, Stephen
 PATENT ASSIGNEE(S): Curis, Inc., USA
 SOURCE: PCT Int. Appl., 123pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008033745	A2	20080320	WO 2007-US77968	20070910
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
US 20080161320	A1	20080703	US 2007-852440	20070910
PRIORITY APPLN. INFO.:			US 2006-843646P	P 20060911
			US 2007-895894P	P 20070320
OTHER SOURCE(S):	MARPAT	148:355814		
GI				

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB Fused bicyclic pyrimidine derivs. I and II [Ar = aryl, substituted arylheteroaryl or heteroaryl; Q = absent or (un)substituted alkyl; X = O, S, NH, or alkylamino; Z = O, S, NR1; Y = N or CR2; B = linker; D = C(O)NH2, NHC(S)CH3, CHC(O)NHacyl, etc.; R1 = H or (un)substituted alkyl; R2 = halo, (un)substituted aliphatic, aryl or heteroaryl], and their pharmaceutically acceptable salts, are prepared and disclosed as protein tyrosine kinase (PTK) inhibitors. Thus, e.g., III was prepared by N-alkylation of 1,4-dioxa-8-azaspiro[4.5]decane with 6-(4-(chloromethyl)phenyl)-N-((R)-1-phenylethyl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine (preparation given) and deprotection followed by condensation with 6-aminohexanoic acid Me ester and amidation with hydroxylamine. Select I were evaluated in EGFR assays, e.g., III demonstrated an IC50 value of ≤ 0.1 μ M.

IT 1011716-90-7P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES

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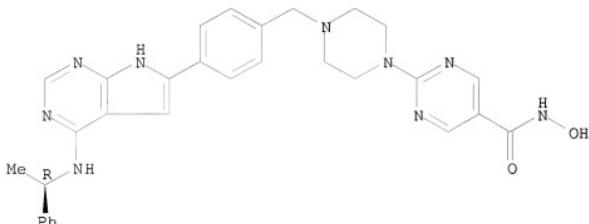
(Uses)

(preparation of (aralkylamino)(phenyl)pyrrolopyrimidine derivs. for use as protein tyrosine kinase (PTK) inhibitors)

RN 1011716-90-7 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[4-[(1R)-1-phenylethyl]amino]-7H-pyrrolo[2,3-d]pyrimidin-6-yl]phenyl]-1-piperazinyl- (CA INDEX NAME)

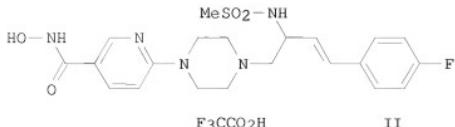
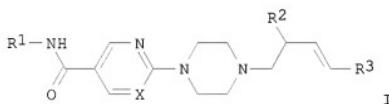
Absolute stereochemistry.



L3 ANSWER 3 OF 11 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2007:816930 CAPLUS
 DOCUMENT NUMBER: 147:211903
 TITLE: Preparation of pyrimidine derivatives as histone deacetylase inhibitors
 INVENTOR(S): Marconnet-Decrane, Laurence Francoise Bernadette; Gaurrand, Sandrine Francoise Dominique; Angibaud, Patrick Rene
 PATENT ASSIGNEE(S): Janssen Pharmaceutica N.V., Belg.
 SOURCE: PCT Int. Appl., 62pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007082874	A1	20070726	WO 2007-EP50371	20070116
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JE, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, IJ, TM				
CA 2630717	A1	20070726	CA 2007-2630717	20070116
PRIORITY APPLN. INFO.:			EP 2006-100570	A 20060119
			WO 2007-EP50371	W 20070116

OTHER SOURCE(S): MARPAT 147:211903
 GI



AB The title compds. with general formula I [wherein R1 = OH or substituted phenyl; X = N or CH; R2 = amino, alkylamino, alkoxy, OH, etc.; R3 = (un)substituted Ph, naphthalene, or heterocycle] or N-oxide forms, pharmaceutically acceptable salts, or stereoisomeric forms thereof were prepared as histone deacetylase (HDAC) inhibitors for the treatment of proliferative diseases. For example, compound II was prepared in a multi-step synthesis. In vitro assay for inhibition of HDAC was performed to measure the inhibition of HDAC enzymic activity, and colorimetric assay was performed to determine cellular activity on A2780 tumor cells. II showed HDAC inhibitory and anti-proliferative activities in the above two assays with pIC50 values of 7.0 and 5.3, resp. Formulations containing I as active ingredients were also reported.

IT 944738-91-4P 944738-94-7P 944738-97-0P

944739-00-8P 944739-08-6P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of pyrimidine derivs. as histone deacetylase inhibitors)

RN 944738-91-4 CAPLUS

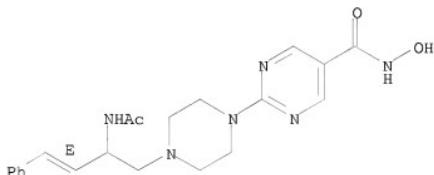
CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(acetylamino)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944738-90-3

CMF C21 H26 N6 O3

Double bond geometry as shown.



CM 2

CRN 76-05-1

CMF C2 H F3 O2



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RN 944738-94-7 CAPLUS

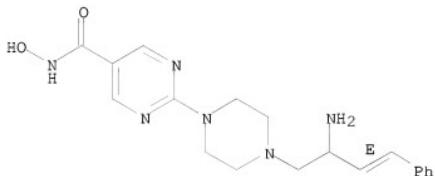
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CM 1

CRN 944738-93-6

CMF C19 H24 N6 O2

Double bond geometry as shown.



CM 2

CRN 76-05-1

CMF C2 H F3 O2



RN 944738-97-0 CAPLUS

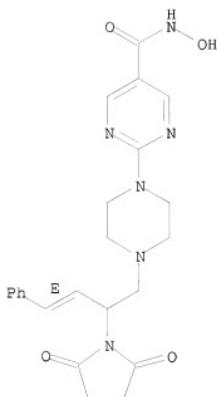
CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(2,5-dioxo-1-pyrrolidinyl)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944738-96-9

CMF C23 H26 N6 O4

Double bond geometry as shown.



CM 2

CRN 76-05-1
CMF C2 H F3 O2

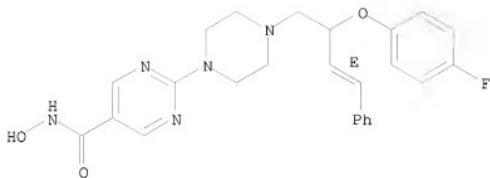
RN 944739-00-8 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(4-fluorophenoxy)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944738-99-2
CMF C25 H26 F N5 O3

Double bond geometry as shown.

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CM 2

CRN 76-05-1
CMF C2 H F3 O2

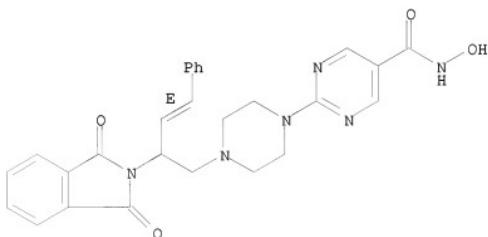


RN 944739-08-6 CAPLUS
CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(1,3-dihydro-1,3-dioxo-2H-isoindol-2-yl)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

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CRN 944739-07-5
CMF C27 H26 N6 O4

Double bond geometry as shown.



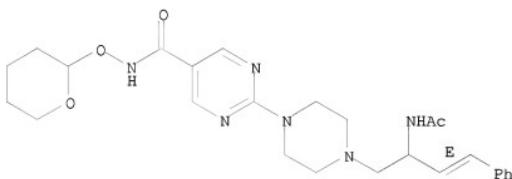
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CRN 76-05-1
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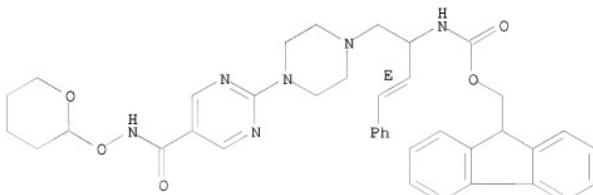
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 944739-36-0P 944739-42-8P 944739-65-5P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
 (Reactant or reagent)
 (preparation of pyrimidine derivs. as histone deacetylase inhibitors)
 RN 944739-19-9 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(acetylamino)-4-phenyl-3-buten-1-yl]-
 1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)

Double bond geometry as shown.



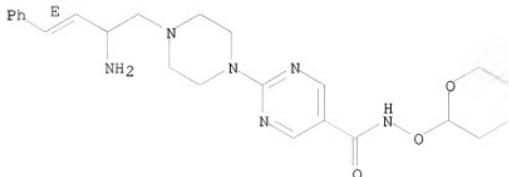
RN 944739-25-7 CAPLUS
 CN Carbamic acid, N-[(2E)-3-phenyl-1-[(4-[(5-[(tetrahydro-2H-pyran-2-yl)oxy]amino)carbonyl]-2-pyrimidinyl]-1-piperazinyl)methyl]-2-propen-1-yl]-, 9H-fluoren-9-ylmethyl ester (CA INDEX NAME)

Double bond geometry as shown.



RN 944739-27-9 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-amino-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)

Double bond geometry as shown.

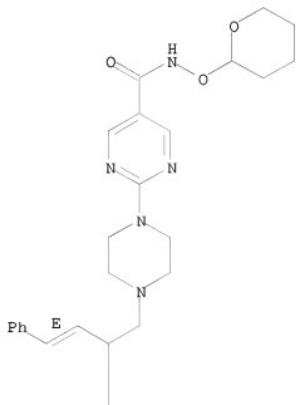


RN 944739-36-0 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(2,5-dioxo-1-pyrrolidinyl)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)

Double bond geometry as shown.

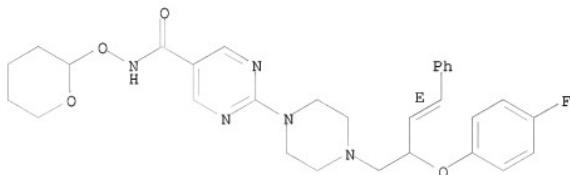
PAGE 1-A





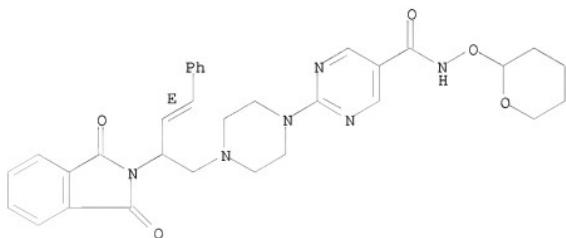
RN 944739-42-8 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(4-fluorophenoxy)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)

Double bond geometry as shown.



RN 944739-65-5 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(1,3-dihydro-1,3-dioxo-2H-isoindol-2-yl)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)

Double bond geometry as shown.



REFERENCE COUNT:

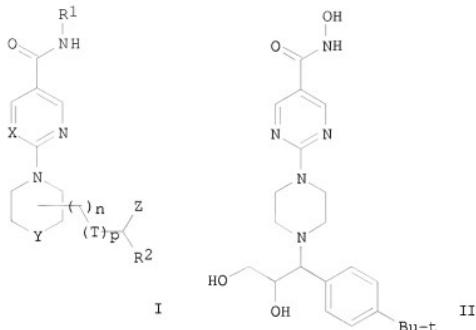
4

THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

10/513699

L3 ANSWER 4 OF 11 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:816806 CAPLUS
DOCUMENT NUMBER: 147:211902
TITLE: Preparation of pyrimidine derivatives as histone deacetylase inhibitors
INVENTOR(S): Angibaud, Patrick Rene; Van Brandt, Sven Franciscus Anna; Marconnet-Decrane, Laurence Francoise Bernadette; Roux, Bruno
PATENT ASSIGNEE(S): Janssen Pharmaceutica N.V., Belg.
SOURCE: PCT Int. Appl., 63pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007082880	A1	20070726	WO 2007-EP50379	20070116
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JE, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, IJ, TM			EP 2006-100571	A 20060119
PRIORITY APPLN. INFO.:				
OTHER SOURCE(S):	MARPAT	147:211902		
GI				



AB The title compds. with general formula I (wherein R1 = OH or substituted phenyl; R2 = -CH2OH, -CH2COH3, -CH2COCH2CH3, or -CH2CH(OH)CH2OH; T = N(R3), where R3 = H, alkyl, cycloalkyl, etc.; X = N or CH; Y = O, NH, CH2, etc.; n = 0-1; p = 0-1, provided that when p = 0 then n = 0 and Y = N, and -CH(R2)-Z is attached to Y; Z = (un)substituted aryl or heteroaryl) or N-oxide forms, pharmaceutically acceptable salts, or stereoisomeric forms thereof were prepared as histone deacetylase (HDAC) inhibitors for the treatment of proliferative diseases. For example, compound II was prepared in a multi-step synthesis. In vitro assay for inhibition of HDAC was performed to measure the inhibition of HDAC enzymic activity, and colorimetric assay was performed to determine cellular activity on A2780 tumor cells. II showed HDAC inhibitory and anti-proliferative activities in the above two assays with pIC50 values of 7.0 and 7.1, resp. Formulations containing I as active ingredients were also reported.

IT 944712-03-2P 944712-05-4P 944712-07-6P

944712-09-8P 944712-10-1P 944712-12-3P

$944712=14=5P$ $944712=16=7P$ $944712=18=9P$

BT: PAC (Pharmacological activity): SPM

RE: PAC (Pharmacological activity); SIR (Synthetic preparation); THER (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of pyrimidine derivs. as histone deacetylase inhibitors)

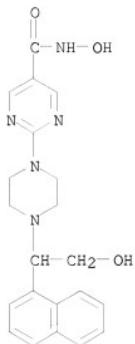
RN 944712-03-2 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-hydroxy-1-(1-naphthalenyl)ethyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-02-1
CMF C21 H23 N5 Q3

10/513699



CM 2

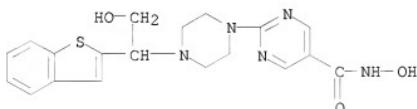
CRN 76-05-1
CMF C2 H F3 O2



RN 944712-05-4 CAPLUS
CN 5-Pyrimidinecarboxamide, 2-[4-(1-benzo[b]thien-2-yl-2-hydroxyethyl)-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-04-3
CMF C19 H21 N5 O3 S



CM 2

10/513699

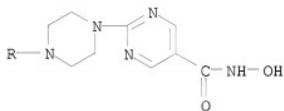
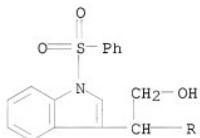
CRN 76-05-1
CMF C2 H F3 O2



RN 944712-07-6 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-hydroxy-1-[1-(phenylsulfonyl)-1H-indol-3-yl]ethyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-06-5
CMF C25 H26 N6 O5 S



CM 2

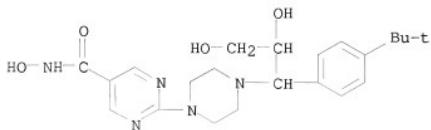
CRN 76-05-1
CMF C2 H F3 O2



RN 944712-09-8 CAPLUS
CN 5-Pyrimidinecarboxamide, 2-[4-[1-[4-(1,1-dimethylethyl)phenyl]-2,3-

10/513699

dihydroxypropyl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



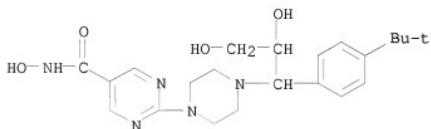
RN 944712-10-1 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1-[4-(1,1-dimethylethyl)phenyl]-2,3-dihydroxypropyl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1)
(CA INDEX NAME)

CM 1

CRN 944712-09-8

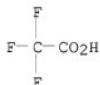
CMF C22 H31 N5 O4



CM 2

CRN 76-05-1

CMF C2 H F3 O2



RN 944712-12-3 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[(1R,2S)-1-[4-(1,1-dimethylethyl)phenyl]-2,3-dihydroxypropyl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?)
(CA INDEX NAME)

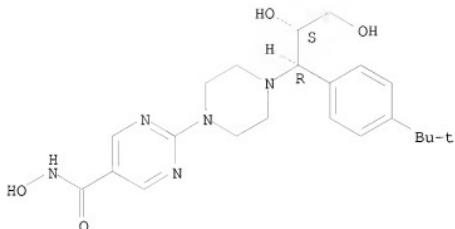
CM 1

CRN 944712-11-2

CMF C22 H31 N5 O4

Absolute stereochemistry.

10/513699



CM 2

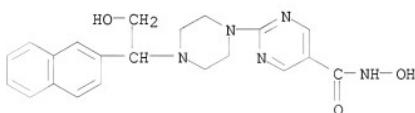
CRN 76-05-1
CMF C2 H F3 O2



RN 944712-14-5 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-hydroxy-1-(2-naphthalenyl)ethyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 944712-13-4
CMF C21 H23 N5 O3



CM 2

CRN 76-05-1
CMF C2 H F3 O2

10/513699

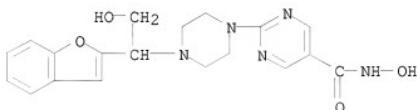


RN 944712-16-7 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1-(2-benzofuranyl)-2-hydroxyethyl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-15-6
CMF C19 H21 N5 O4



CM 2

CRN 76-05-1
CMF C2 H F3 O2

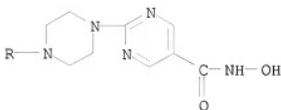
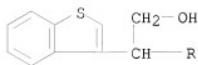


RN 944712-18-9 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-(1-benzo[b]thien-3-yl-2-hydroxyethyl)-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-17-8
CMF C19 H21 N5 O3 S



CM 2

CRN 76-05-1
CMF C2 H F3 O2

IT 944712-19-0P 944712-20-3P 944712-23-6P

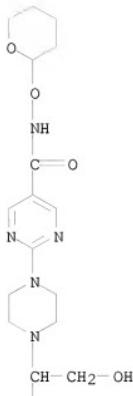
944712-27-0P 944712-30-5P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

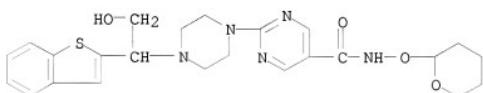
(preparation of pyrimidine derivs. as histone deacetylase inhibitors)

RN 944712-19-0 CAPLUS

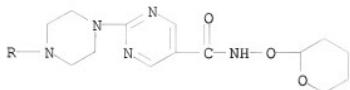
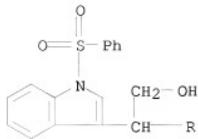
CN 5-Pyrimidinecarboxamide, 2-[4-[2-hydroxy-1-(1-naphthalenyl)ethyl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)



RN 944712-20-3 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-(1-benzo[b]thien-2-yl-2-hydroxyethyl)-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)

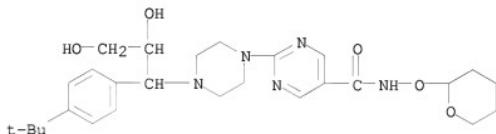


RN 944712-23-6 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[2-hydroxy-1-[1-(phenylsulfonyl)-1H-indol-3-yl]ethyl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)



RN 944712-27-0 CAPLUS

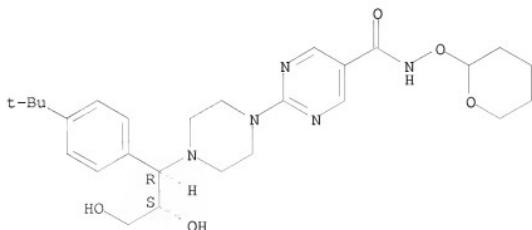
CN 5-Pyrimidinecarboxamide, 2-[4-[1-[4-(1,1-dimethylethyl)phenyl]-2,3-dihydroxypropyl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)



RN 944712-30-5 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[(1R,2S)-1-[4-(1,1-dimethylethyl)phenyl]-2,3-dihydroxypropyl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)

Absolute stereochemistry.



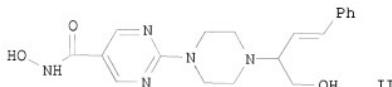
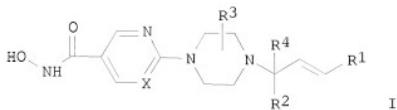
10/513699

REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 5 OF 11 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2006:101446 CAPLUS
 DOCUMENT NUMBER: 144:192266
 TITLE: Preparation of substituted propenyl piperazine derivatives as novel inhibitors of histone deacetylase
 INVENTOR(S): Van Brandt, Sven Franciscus Anna; Van Emelen, Kristof;
 Angibaud, Patrick Rene; Marconnet-Decrane, Laurence
 Francoise Bernadette; Arts, Janine
 PATENT ASSIGNEE(S): Janssen Pharmaceutica N.V., Belg.
 SOURCE: PCT Int. Appl., 67 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006010749	A2	20060202	WO 2005-EP53611	20050725
WO 2006010749	A3	20060608		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
AU 2005266311	A1	20060202	AU 2005-266311	20050725
CA 2572971	A1	20060202	CA 2005-2572971	20050725
EP 1776358	A2	20070425	EP 2005-777776	20050725
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, YU				
CN 1993356	A	20070704	CN 2005-80025487	20050725
JP 2008508234	T	20080321	JP 2007-523072	20050725
BR 2005013891	A	20080520	BR 2005-13891	20050725
KR 2007043978	A	20070426	KR 2007-701641	20070123
US 20070135424	A1	20070614	US 2007-626215	20070123
IN 2007DN00658	A	20070803	IN 2007-DN658	20070124
MX 200701119	A	20070315	MX 2007-1119	20070126
NO 200700117	A	20070227	NO 2007-1117	20070227
PRIORITY APPLN. INFO.:			EP 2004-77171	A 20040728
			US 2004-592357P	P 20040729
			WO 2005-EP53611	W 20050725

OTHER SOURCE(S): CASREACT 144:192266; MARPAT 144:192266
 GI



AB Substituted propenyl piperazine derivs. I, wherein X is independently N or CH; R1 is Ph, naphthalenyl or heterocyclyl; wherein each of said Ph or naphthalenyl is optionally substituted with one or two substituents each independently selected from halo, alkyl, alkyloxy, poly-halo-alkyl, aryl, hydroxy, cyano, amino, alkylcarbonylamino, alkylsulfonylamino, hydroxycarbonyl, alkyloxycarbonyl, hydroxyalkyl, alkyloxymethyl, aminomethyl, alkylaminomethyl, alkylcarbonylaminomethyl, alkylsulfonylaminomethyl, aminosulfonyl, alkylaminosulfonyl or heterocyclyl; R2 is hydrogen, -CH2R5, trifluoromethyl, -C(O)-R6, or -CH-NR/R8; wherein each R5 is independently hydrogen, hydroxy, alkyloxy, alkyloxyalkyloxy, alkylcarbonyloxy, piperazinyl, N-methylpiperazinyl, morpholinyl, thiomorpholinyl, imidazolyl or triazolyl; each R6 is independently hydroxy, alkyloxy, amino or mono- or di(alkyl)amino, cycloalkylamino, hydroxylalkylamino, piperazinyl, N-methylpiperazinyl, morpholinyl or thiomorpholinyl; each R7 and R8 are independently hydrogen, alkyl, alkylcarbonyl, alkylsulfonyl, or mono- or di(alkyl)aminosulfonyl; R3 is hydrogen, hydroxymethyl, aminomethyl or mono- or di(alkyl)aminomethyl; R4 is hydrogen or alkyl; were prepared and having histone deacetylase inhibiting enzymic activity and to inhibit proliferative conditions, such as cancer and psoriasis. Thus, propenyl piperazine derivative II was prepared and tested in vitro and in nude mice as inhibitor of histone deacetylase and was better than R306465 after oral administration. P21 enzyme linked immunosorbent assay has been applied to determine the p21 protein expression level in human A2780 ovarian carcinoma cells. In vitro assay for inhibition of histone deacetylase is reported. P21 induction was measured as the consequence of DNA damage or as the consequence of histone deacetylase inhibition. Antiproliferative activity of title compds. was determined on A2780 cells (neg. log value of the IC50, pIC50 = 7.9-8.2).

IT

875138-85-5P 875138-87-7P 875138-88-8P
 875138-89-9P 875138-90-2P 875138-91-3P
 875138-93-5P 875138-94-6P 875138-98-0P
 875139-00-7P 875139-02-9P 875139-04-1P
 875139-06-3P 875139-07-4P 875139-09-6P
 875139-11-0P 875139-13-2P 875139-14-3P
 875139-15-4P 875139-17-6P 875139-19-8P
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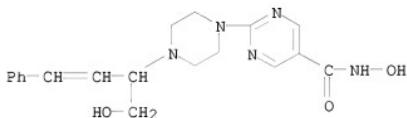
875139-70-1P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of substituted propenyl piperazine derivs. as novel inhibitors of histone deacetylase)

RN 875138-85-5 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(1-(hydroxymethyl)-3-phenyl-2-propen-1-yl)-1-piperazinyl]- (CA INDEX NAME)



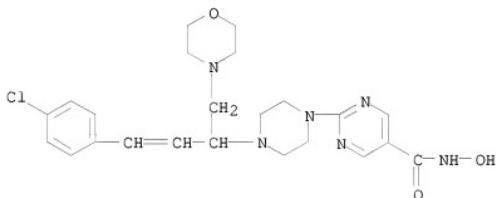
RN 875138-87-7 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3-(4-chlorophenyl)-1-(4-morpholinylmethyl)-2-propen-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-86-6

CMF C23 H29 Cl N6 O3



CM 2

CRN 76-05-1

CMF C2 H F3 O2

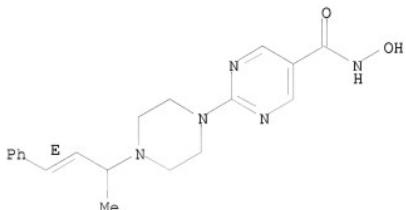


10/513699

RN 875138-88-8 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-methyl-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)

Double bond geometry as shown.



RN 875138-89-9 CAPLUS

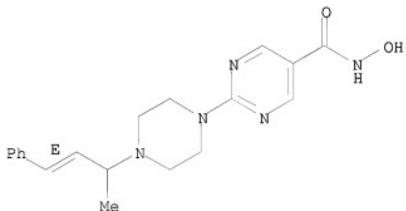
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-methyl-3-phenyl-2-propen-1-yl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-88-8

CMF C19 H23 N5 O2

Double bond geometry as shown.



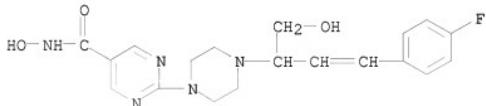
CM 2

CRN 76-05-1

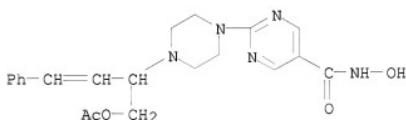
CMF C2 H F3 O2



RN 875138-90-2 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[3-(4-fluorophenyl)-1-(hydroxymethyl)-2-propen-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

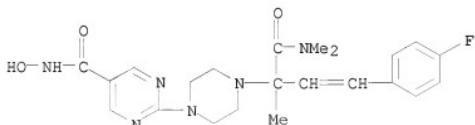


RN 875138-91-3 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(acetoxy)methyl]-3-phenyl-2-propen-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



RN 875138-93-5 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(dimethylamino)carbonyl]-3-(4-fluorophenyl)-1-methyl-2-propen-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-92-4
CMF C22 H27 F N6 O3

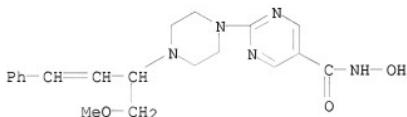
CM 2

10/513699

CRN 76-05-1
CMF C2 H F3 O2



RN 875138-94-6 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-(methoxymethyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)

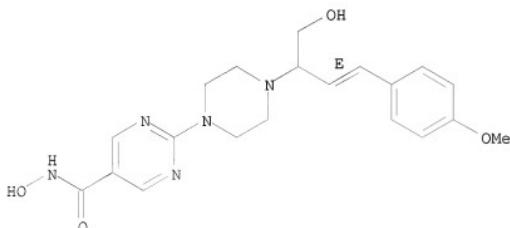


RN 875138-98-0 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-(hydroxymethyl)-3-(4-methoxyphenyl)-2-propen-1-yl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-97-9
CMF C20 H25 N5 O4

Double bond geometry as shown.



CM 2

CRN 76-05-1

10/513699

CMF C2 H F3 O2



RN 875139-00-7 CAPLUS

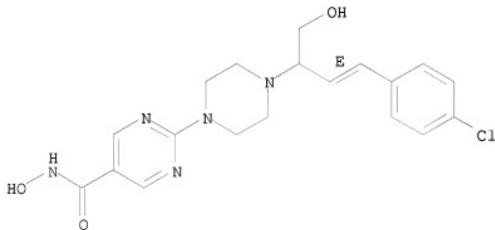
CN 5-Pyrimidinecarboxamide, 2-[4-[(2E)-3-(4-chlorophenyl)-1-(hydroxymethyl)-2-propen-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-99-1

CMF C19 H22 Cl N5 O3

Double bond geometry as shown.



CM 2

CRN 76-05-1

CMF C2 H F3 O2



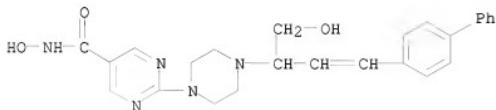
RN 875139-02-9 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3-[1,1'-biphenyl]-4-yl]-1-(hydroxymethyl)-2-propen-1-yl]-1-piperazinyl-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

10/513699

CRN 875139-01-8
CMF C25 H27 N5 O3



CM 2

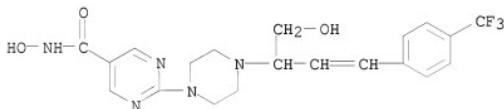
CRN 76-05-1
CMF C2 H F3 O2



RN 875139-04-1 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-(hydroxymethyl)-3-[4-(trifluoromethyl)phenyl]-2-propenyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-03-0
CMF C20 H22 F3 N5 O3



CM 2

CRN 76-05-1
CMF C2 H F3 O2

10/513699



RN 875139-06-3 CAPLUS

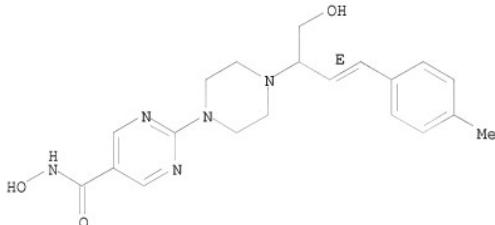
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-(hydroxymethyl)-3-(4-methylphenyl)-2-propen-1-yl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-05-2

CMF C20 H25 N5 O3

Double bond geometry as shown.



CM 2

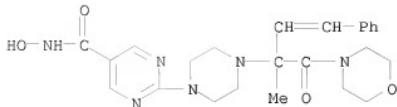
CRN 76-05-1

CMF C2 H F3 O2



RN 875139-07-4 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-methyl-1-(4-morpholinylcarbonyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)



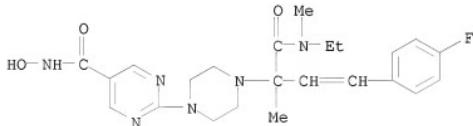
RN 875139-09-6 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(ethylmethylamino)carbonyl]-3-(4-fluorophenyl)-1-methyl-2-propenyl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-08-5

CMF C23 H29 F N6 O3



CM 2

CRN 76-05-1

CMF C2 H F3 O2



RN 875139-11-0 CAPLUS

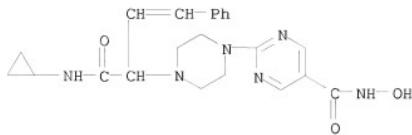
CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(cyclopropylamino)carbonyl]-3-phenyl-2-propenyl-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-10-9

CMF C22 H26 N6 O3

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2

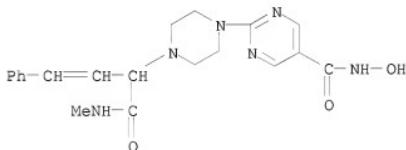


RN 875139-13-2 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(1-[(methylamino)carbonyl]-3-phenyl-2-propen-1-yl)-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-12-1
CMF C20 H24 N6 O3



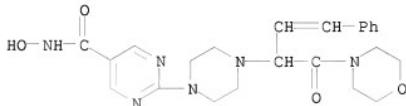
CM 2

CRN 76-05-1
CMF C2 H F3 O2



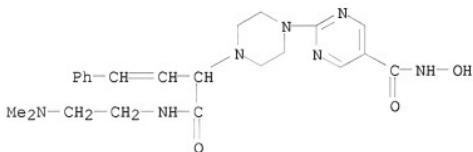
RN 875139-14-3 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-(4-morpholinylcarbonyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)



RN 875139-15-4 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(2-(dimethylamino)ethyl]amino]carbonyl]-3-phenyl-2-propen-1-yl]-1-piperazinyl-N-hydroxy- (CA INDEX NAME)



RN 875139-17-6 CAPLUS

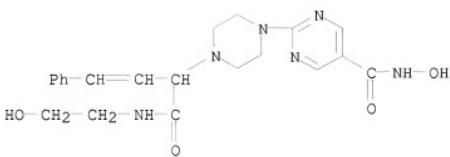
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-[(2-hydroxyethyl)amino]carbonyl]-3-phenyl-2-propen-1-yl]-1-piperazinyl-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-16-5

CMF C21 H26 N6 O4

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2

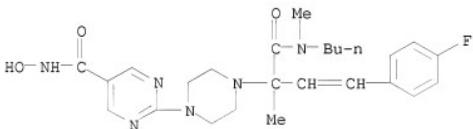


RN 875139-19-8 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[(butylmethylamino)carbonyl]-3-(4-fluorophenyl)-2-propenyl-1-yl]-1-piperazinyl-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-18-7
CMF C25 H33 F N6 O3

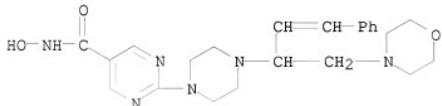


CM 2

CRN 76-05-1
CMF C2 H F3 O2

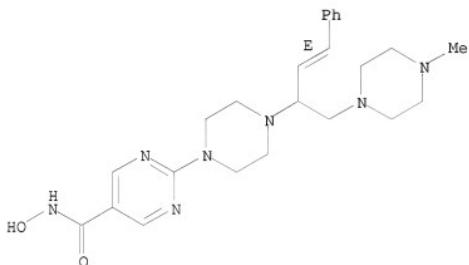


RN 875139-20-1 CAPLUS
 CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-(4-morpholinylmethyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)



RN 875139-21-2 CAPLUS
 CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-[(4-methyl-1-piperazinyl)methyl]-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)

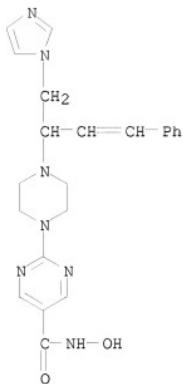
Double bond geometry as shown.



RN 875139-23-4 CAPLUS
 CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-(1H-imidazol-1-ylmethyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

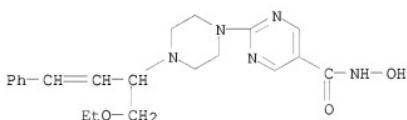
CRN 875139-22-3
 CMF C22 H25 N7 O2



CM 2

CRN 76-05-1
CMF C2 H F3 O2

RN 875139-24-5 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-(1-(ethoxymethyl)-3-phenyl-2-propen-1-yl)-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

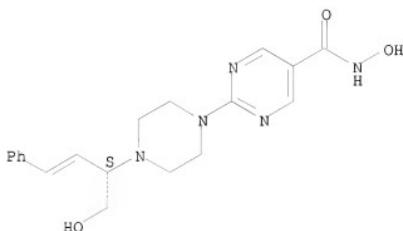


RN 875139-25-6 CAPLUS
 CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(1S)-1-(hydroxymethyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)

Absolute stereochemistry.

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Double bond geometry unknown.

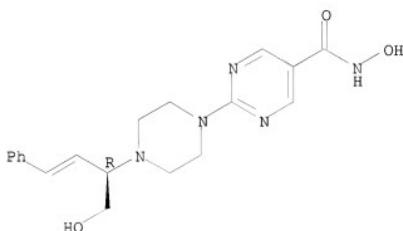


RN 875139-26-7 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(1R)-1-(hydroxymethyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry unknown.

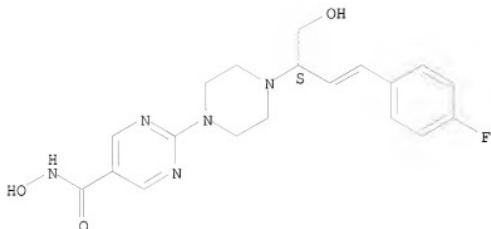


RN 875139-27-8 CAPLUS

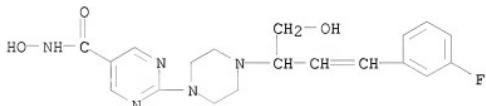
CN 5-Pyrimidinecarboxamide, 2-[4-[(1S)-3-(4-fluorophenyl)-1-(hydroxymethyl)-2-propen-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

Absolute stereochemistry.

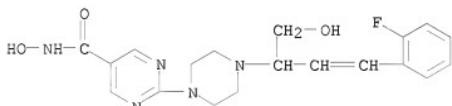
Double bond geometry unknown.



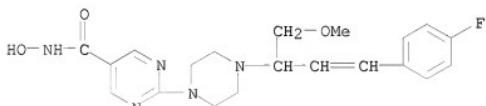
RN 875139-28-9 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[3-(3-fluorophenyl)-1-(hydroxymethyl)-2-propenyl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



RN 875139-29-0 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[3-(2-fluorophenyl)-1-(hydroxymethyl)-2-propenyl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



RN 875139-30-3 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[3-(4-fluorophenyl)-1-(methoxymethyl)-2-propenyl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

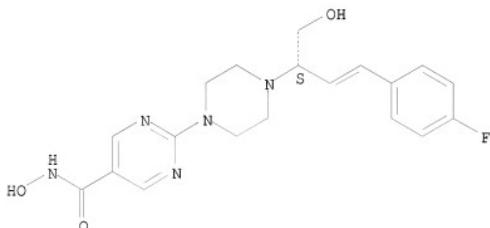


RN 875139-31-4 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[(1S)-3-(4-fluorophenyl)-1-(hydroxymethyl)-2-

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propen-1-yl]-1-piperazinyl]-N-hydroxy-, hydrochloride (1:1) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.

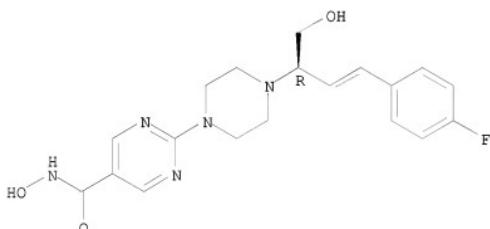


● HCl

RN 875139-69-8 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-((1R)-3-(4-fluorophenyl)-1-(hydroxymethyl)-2-propen-1-yl)-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

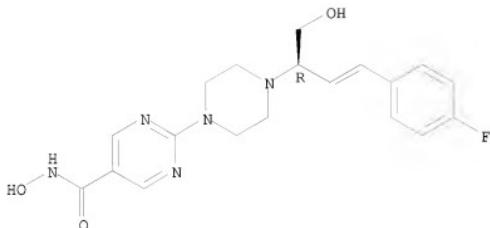
Absolute stereochemistry.
Double bond geometry unknown.



RN 875139-70-1 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-((1R)-3-(4-fluorophenyl)-1-(hydroxymethyl)-2-propen-1-yl)-1-piperazinyl]-N-hydroxy-, hydrochloride (1:1) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.



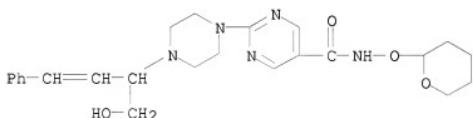
● HCl

IT 875138-54-8P 875138-59-3P 875138-62-8P
 875138-66-2P 875138-70-8P 875138-73-1P
 875138-77-5P 875138-78-6P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (preparation of substituted propenyl piperazine derivs. as novel inhibitors of histone deacetylase)

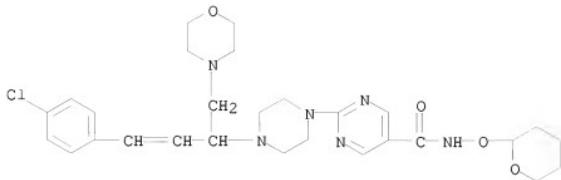
RN 875138-54-8 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1-(hydroxymethyl)-3-phenyl-2-propenyl-1-yl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)



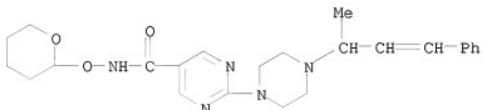
RN 875138-59-3 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3-(4-chlorophenyl)-1-(4-morpholinylmethyl)-2-propenyl-1-yl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)



RN 875138-62-8 CAPLUS

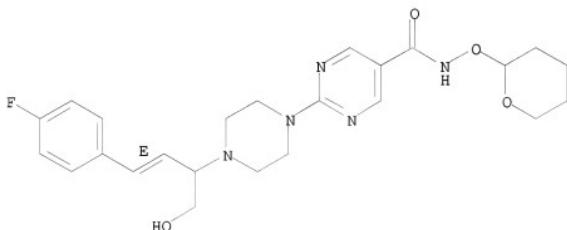
CN 5-Pyrimidinecarboxamide, 2-[4-(1-methyl-3-phenyl-2-propen-1-yl)-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)



RN 875138-66-2 CAPLUS

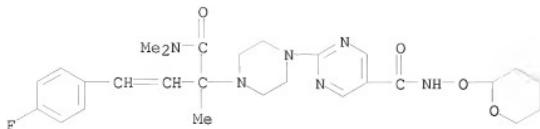
CN 5-Pyrimidinecarboxamide, 2-[4-[(2E)-3-(4-fluorophenyl)-1-(hydroxymethyl)-2-propen-1-yl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)

Double bond geometry as shown.

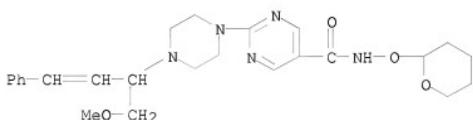


RN 875138-70-8 CAPLUS

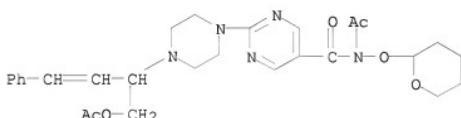
CN 5-Pyrimidinecarboxamide, 2-[4-[1-((dimethylamino)carbonyl)-3-(4-fluorophenyl)-1-methyl-2-propen-1-yl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)



RN 875138-73-1 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[1-(methoxymethyl)-3-phenyl-2-propenyl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)



RN 875138-77-5 CAPLUS
 CN 5-Pyrimidinecarboxamide, N-acetyl-2-[4-[1-[(acetyloxy)methyl]-3-phenyl-2-propenyl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)

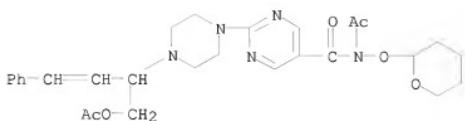


RN 875138-78-6 CAPLUS
 CN 5-Pyrimidinecarboxamide, N-acetyl-2-[4-[1-[(acetoxy)methyl]-3-phenyl-2-propenyl]-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]-, ethanedioate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-77-5
 CMF C28 H35 N5 O6

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CM 2

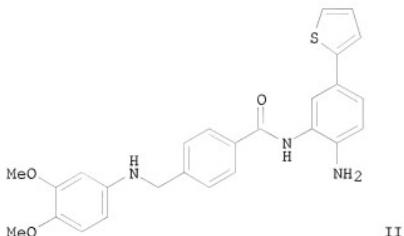
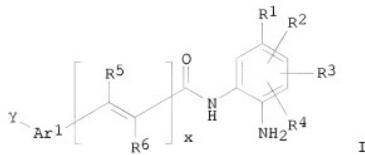
CRN 144-62-7
CMF C2 H2 O4



L3 ANSWER 6 OF 11 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2005:300395 CAPLUS
 DOCUMENT NUMBER: 142:355054
 TITLE: Preparation of amide derivatives as inhibitors of histone deacetylase
 INVENTOR(S): Moradei, Oscar; Paquin, Isabelle; Leit, Silvana;
 Frechette, Sylvie; Vaisburg, Arkadii; Besterman,
 Jeffrey M.; Tessier, Pierre; Mallais, Tammy C.
 PATENT ASSIGNEE(S): Methygene, Inc., Can.
 SOURCE: PCT Int. Appl., 559 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005030705	A1	20050407	WO 2004-US31591	20040924
WO 2005030705	A9	20060420		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2004276337	A1	20050407	AU 2004-276337	20040924
CA 2539117	A1	20050407	CA 2004-2539117	20040924
EP 1663953	A1	20060607	EP 2004-789074	20040924
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR CN 1882529	A	20061220	CN 2004-80034571	20040924
JP 2007506785	T	20070322	JP 2006-528279	20040924
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			US 2003-532973P	P 20031229
			US 2004-561082P	P 20040409
			JP 2006-528279	A3 20040924
			WO 2004-US31591	W 20040924

OTHER SOURCE(S): CASREACT 142:355054; MARPAT 142:355054
 GI



AB Title compds. I [Ar1 = (un)saturated-, (un)substituted-mono or fused poly-cyclic hydrocarbyl optionally containing 1-4 heteroatoms per ring; R1 = (un)substituted-mono-, -bi-, -tri-cyclic-aryl or -heteroaryl; R2, R3, and R4 independently = H, halo, amino, etc.; R5 and R6 independently = H, alkyl, aryl, etc.; x = 0-1; Y = any pharmaceutically acceptable chemical moiety consisting of 1 to 50 atoms with provisions] and their pharmaceutically acceptable salts, are prepared and disclosed as inhibitors of histone deacetylase. Thus, e.g., II was prepared by Suzuki coupling of 2-bromo-2-nitro-phenylamine (preparation given) with 2-thiopheneboronic acid followed by carbonylation with 4-[3,4-dimethoxy-(phenylamino)-methyl]benzoic acid (preparation given) and subsequent reduction. The inhibitory capability of I towards antiproliferative activity of histone deacetylase enzyme was evaluated using 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT) assay and it revealed that certain compds. of the invention had MTT IC 50 values in the range of below 1 up to 20 μ M. I as histone deacetylase inhibitors should prove useful in the treatment of diseases such as, but not limited to, cell proliferative disease, protozoal disease, and fungal disease.

IT 603985-86-0P 603985-88-2P 603985-90-6P
 603985-94-0P 603991-95-3P 603991-96-4P
 603992-24-1P 603992-25-2P 603992-26-3P
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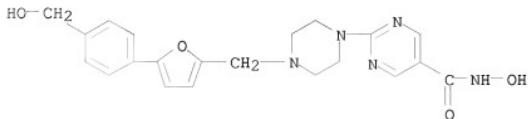
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of amide derivs. as inhibitors of histone deacetylase)

RN 603985-86-0 CAPLUS

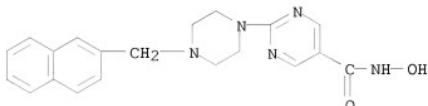
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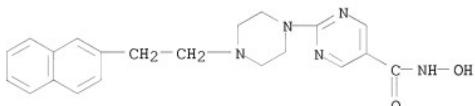
RN 603985-88-2 CAPLUS

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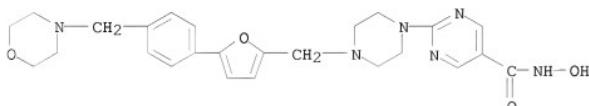
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RN 603985-94-0 CAPLUS

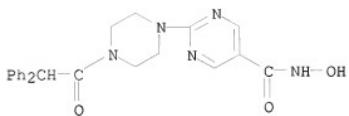
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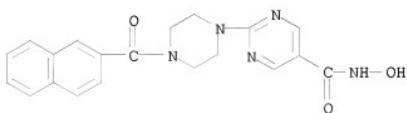
RN 603991-95-3 CAPLUS

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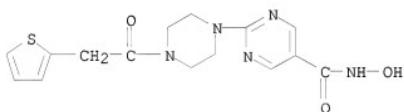
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RN 603991-96-4 CAPLUS
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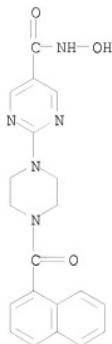


RN 603992-24-1 CAPLUS
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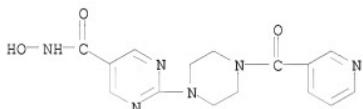


RN 603992-25-2 CAPLUS
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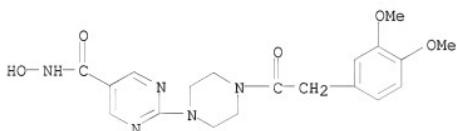
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RN 603992-26-3 CAPLUS
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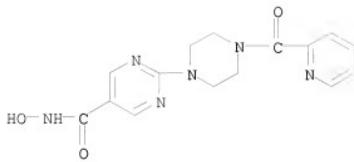


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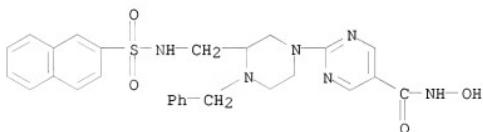
RN 603992-28-5 CAPLUS
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10/513699



RN 604784-81-8 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[3-[(2-naphthalenylsulfonyl)amino]methoxy]-4-(phenylmethyl)-1-piperazinyl- (CA INDEX NAME)



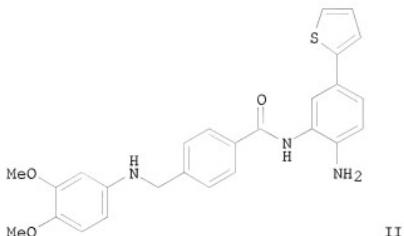
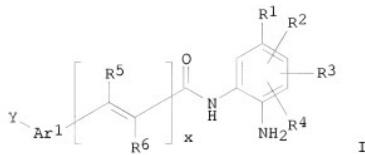
REFERENCE COUNT:

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THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 7 OF 11 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2005:300394 CAPLUS
 DOCUMENT NUMBER: 142:373563
 TITLE: Preparation of amide derivatives as inhibitors of histone deacetylase
 INVENTOR(S): Moradei, Oscar; Paquin, Isabelle; Leit, Silvana;
 Frechette, Sylvie; Vaisburg, Arkadii; Besterman,
 Jeffrey M.; Tessier, Pierre; Mallais, Tammy C.
 PATENT ASSIGNEE(S): Methylgene, Inc., Can.
 SOURCE: PCT Int. Appl., 389 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005030704	A1	20050407	WO 2004-US31590	20040924
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, T2, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
JP 2008094847	A	20080424	JP 2007-281356 US 2003-505884P US 2003-532973P US 2004-561082P JP 2006-528279	20071030 P 20030924 P 20031229 P 20040409 A3 20040924
PRIORITY APPLN. INFO.:				
OTHER SOURCE(S): GI	CASREACT 142:373563; MARPAT 142:373563			



AB Title compds. I [Ar1 = (un)saturated-, (un)substituted-mono or fused poly-cyclic hydrocarbyl optionally containing 1-4 heteroatoms per ring; R1 = (un)substituted-mono-, -bi-, -tri-cyclic-aryl or -heteroaryl; R2, R3, and R4 independently = H, halo, amino, etc.; R5 and R6 independently = H, alkyl, aryl, etc.; x = 0-1; Y = any pharmaceutically acceptable chemical moiety consisting of 1 to 50 atoms with provisions] and their pharmaceutically acceptable salts, are prepared and disclosed as inhibitors of histone deacetylase. Thus, e.g., II was prepared by Suzuki coupling of 2-bromo-2-nitro-phenylamine (preparation given) with 2-thiopheneboronic acid followed by carbonylation with 4-[3,4-dimethoxy-(phenylamino)-methyl]benzoic acid (preparation given) and subsequent reduction. The inhibitory capability of I towards antiproliferative activity of histone deacetylase enzyme was evaluated using 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT) assay and it revealed that certain compds. of the invention had MTT IC 50 values in the range of below 1 up to 20 μ M. I as histone deacetylase inhibitors should prove useful in the treatment of diseases such as, but not limited to, cell proliferative disease, protozoal disease, and fungal disease.

IT 603985-86-0P 603985-88-2P 603985-90-6P
 603985-94-0P 603991-95-3P 603991-96-4P
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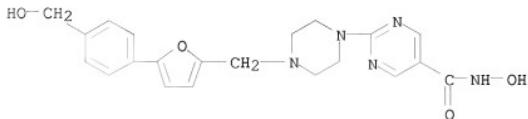
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of amide derivs. as inhibitors of histone deacetylase)

RN 603985-86-0 CAPLUS

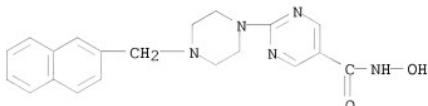
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CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(5-[4-(hydroxymethyl)phenyl]-2-furanyl)methyl]-1-piperazinyl]- (CA INDEX NAME)



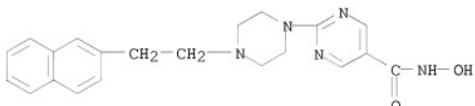
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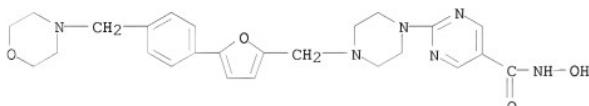
RN 603985-90-6 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-(2-naphthalenyl)ethyl]-1-piperazinyl]- (CA INDEX NAME)



RN 603985-94-0 CAPLUS

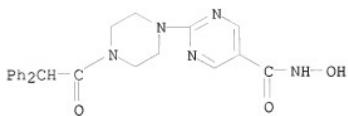
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(5-[4-(4-morpholinylmethyl)phenyl]-2-furanyl)methyl]-1-piperazinyl]- (CA INDEX NAME)



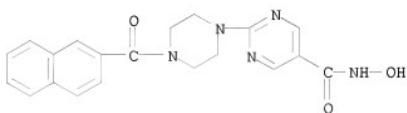
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CN 5-Pyrimidinecarboxamide, 2-[4-(2,2-diphenylacetyl)-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

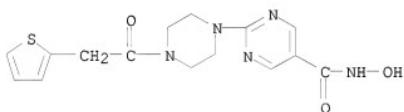
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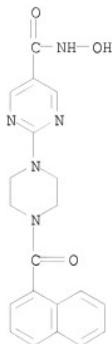
RN 603991-96-4 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-naphthalenylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)



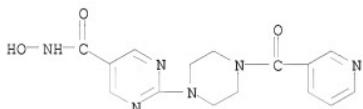
RN 603992-24-1 CAPLUS
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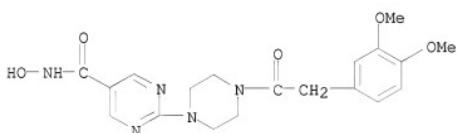
RN 603992-25-2 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(1-naphthalenylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)



RN 603992-26-3 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(3-pyridinylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)

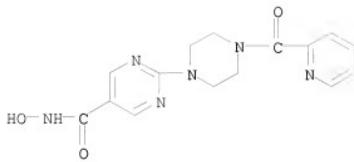


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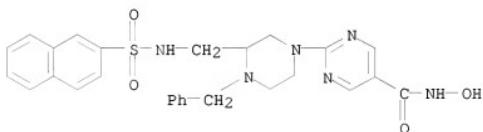
RN 603992-28-5 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-pyridinylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)

10/513699



RN 604784-81-8 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[3-[(2-naphthalenylsulfonyl)amino]methoxy]-4-(phenylmethyl)-1-piperazinyl- (CA INDEX NAME)



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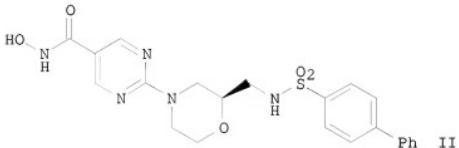
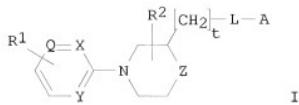
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THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 8 OF 11 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2003:737757 CAPLUS
 DOCUMENT NUMBER: 139:276911
 TITLE: Preparation of N-(piperazinylmethyl-, piperidinylmethyl- and morpholinylmethyl) sulfonamides and amides as novel inhibitors of histone deacetylase
 INVENTOR(S): Van Emelen, Kristof
 PATENT ASSIGNEE(S): Janssen Pharmaceutica N.V., Belg.
 SOURCE: PCT Int. Appl., 69 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 8
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003076438	A1	20030918	WO 2003-EP2510	20030311
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KE, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
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EP 1485378	A1	20041215	EP 2003-711979	20030311
EP 1485378	B1	20080618		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
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CN 1642948	A	20050720	CN 2003-805921	20030311
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NZ 534833	A	20060728	NZ 2003-534833	20030311
CN 101007803	A	20070801	CN 2007-10005212	20030311
AT 398615	T	20080715	AT 2003-711979	20030311
TW 283676	B	20070711	TW 2003-92105285	20030312
IN 2004DN02536	A	20070413	IN 2004-DN2536	20040831
US 20050165016	A1	20050728	US 2004-507084	20040908
MX 2004PA08795	A	20041126	MX 2004-PA8795	20040910
NO 2004004135	A	20040929	NO 2004-4135	20040929
PRIORITY APPLN. INFO.:			US 2002-363799P	P 20020313
			WO 2002-EP14833	A 20021223
			CN 2003-805921	A3 20030311
			WO 2003-EP2510	W 20030311

OTHER SOURCE(S): MARPAT 139:276911
 GI

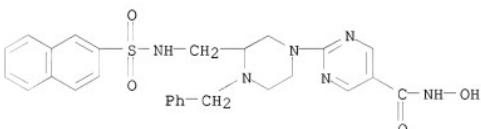


AB The title compds. [I; t = 0-4; Q, X, Y = N, C; Z = NH, O, CH₂; R1 = CONR₃R₄, NHCOR₇, CO(alkanediyl)SR₇, etc. (wherein R₃, R₄ = H, OH, alkyl, etc.; R₇ = H, alkyl, alkylcarbonyl, etc.); R2 = H, OH, NH₂, etc.; L = NR₉CO, NR₉SO₂, NR₉CH₂ (R₉ = H, alkyl, cycloalkyl, etc.); A = (un)substituted Ph, cycloalkyl, pyridyl, etc.] , having histone deacetylase inhibiting enzymic activity, were prepared and formulated. E.g., a multi-step synthesis of (+)-II which showed pIC₅₀ of 7.723 against HDAC, was given.

IT 604784-81-8P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of N-(piperazinylmethyl-, piperidinylmethyl- and morpholinylmethyl) sulfonamides and amides as novel inhibitors of histone deacetylase)



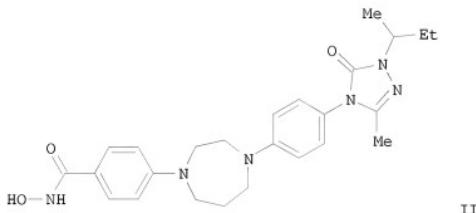
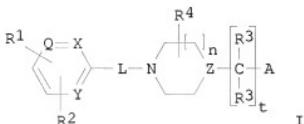
REFERENCE COUNT:

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THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE REFORMAT.

L3 ANSWER 9 OF 11 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2003:737723 CAPLUS
 DOCUMENT NUMBER: 139:261309
 TITLE: Preparation of N-hydroxy-5-piperazino(piperidino or diazepino)-2-pyrimidinecarboxamides and N-hydroxy-4-piperazino(piperidino or diazepino)benzamides as new inhibitors of histone deacetylase
 INVENTOR(S): Angibaud, Patrick Rene; Pilatte, Isabelle Noelle Constance; Van Brandt, Sven Franciscus Anna; Roux, Bruno; Ten Holte, Peter; Verdonck, Marc Gustaaf Celine; Meerpoel, Lieven; Dyatkin, Alexey Borisovich Janssen Pharmaceutica N.V., Belg.
 PATENT ASSIGNEE(S):
 SOURCE: PCT Int. Appl., 72 pp.
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 8
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003076400	A1	20030918	WO 2003-EP2514	20030311
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
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AU 2003218736	A1	20030922	AU 2003-218736	20030311
EP 1485353	A1	20041215	EP 2003-711980	20030311
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BR 2003008081	A	20041221	BR 2003-8081	20030311
CN 1639125	A	20050713	CN 2003-805675	20030311
CN 1642551	A	20050720	CN 2003-805833	20030311
NZ 534834	A	20050729	NZ 2003-534834	20030311
JP 2005526067	T	20050902	JP 2003-574621	20030311
CN 101007803	A	20070801	CN 2007-10005212	20030311
IN 2004DN02533	A	20070413	IN 2004-DN2533	20040831
US 20050107384	A1	20050519	US 2004-506998	20040908
ZA 2004007237	A	20050928	ZA 2004-7237	20040909
ZA 2004007235	A	20051004	ZA 2004-7235	20040909
ZA 2004007232	A	20051006	ZA 2004-7232	20040909
ZA 2004007233	A	20051006	ZA 2004-7233	20040909
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ZA 2004007236	A	20051006	ZA 2004-7236	20040909
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NO 2004004194	A	20041001	NO 2004-4194	20041001
PRIORITY APPLN. INFO.:				
		US 2002-363799P	P	20020313
		WO 2002-EP14833	A	20021223
		CN 2003-805921	A3	20030311



AB The title compds. [I; n = 0-3; t = 0-4; Q, X, Y = N, C; Z = N, CH; R1 = CONR7R8, NHCOR9, CO(alkanediyl)SR9, etc. (wherein R7, R8 = H, OH, alkyl, etc.; R9 = H, alkyl, alkylcarbonyl, etc.); R2 = H, halo, OH, etc.; L = a bond, alkanediyl, alkanediyoxy, NH, CO, NHCO; each R3 = H and one H atom can be replaced by aryl; R4 = H, OH, NH2, etc.; A = (un)substituted Ph, cyclohexyl, pyridyl, etc.], having histone deacetylase inhibiting enzymic activity, were prepared and formulated. E.g., a multi-step synthesis of II which showed pIC50 of 5.121 against HDAC, was given.

IT 603985-87-1P 603985-89-3P 603985-91-7P

603985-95-1P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of piperazino(piperidino or diazepino) substituted 2-pyrimidinecarbohydroxamic acids and N-hydroxybenzamides as new inhibitors of histone deacetylase)

RN 603985-87-1 CAPLUS

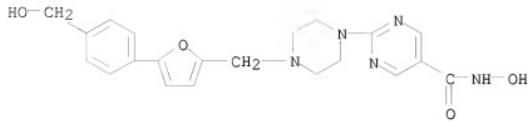
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-([5-[4-(hydroxymethyl)phenyl]-2-furanyl)methyl]-1-piperazinyl-, 2,2,2-trifluoroacetate (5:4) (CA INDEX NAME)

CM 1

CRN 603985-86-0

CMF C21 H23 N5 O4

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2

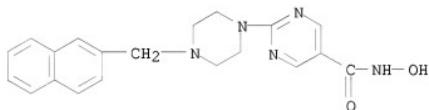


RN 603985-89-3 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-naphthalenylmethyl)-1-piperazinyl]-, 2,2,2-trifluoroacetate (5:4) (CA INDEX NAME)

CM 1

CRN 603985-88-2
CMF C20 H21 N5 O2



CM 2

CRN 76-05-1
CMF C2 H F3 O2



RN 603985-91-7 CAPLUS

<12/04/2007>

Erich Leese

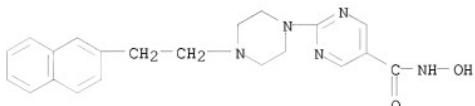
10/513699

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-(2-naphthalenyl)ethyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (5:4) (CA INDEX NAME)

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CRN 603985-90-6

CMF C21 H23 N5 O2



CM 2

CRN 76-05-1

CMF C2 H F3 O2



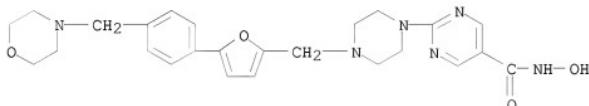
RN 603985-95-1 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[5-[4-(4-morpholinylmethyl)phenyl]-2-furanyl]methyl]-1-piperazinyl-, 2,2,2-trifluoroacetate (1:2) (CA INDEX NAME)

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CRN 603985-94-0

CMF C25 H30 N6 O4



CM 2

CRN 76-05-1

CMF C2 H F3 O2



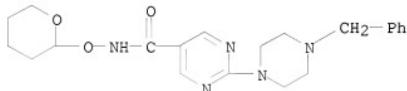
IT 603986-73-8P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of piperazino(piperidino or diazepino) substituted 2-pyrimidinecarboxylic acids and N-hydroxybenzamides as new inhibitors of histone deacetylase)

RN 603986-73-8 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-(phenylmethyl)-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)



REFERENCE COUNT:

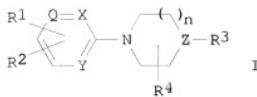
3

THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 10 OF 11 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2003:737586 CAPLUS
 DOCUMENT NUMBER: 139:261308
 TITLE: Preparation of aryl and heteroaryl hydroxamic acids as inhibitors of histone deacetylase for treating proliferative diseases
 INVENTOR(S): Van Emelen, Kristof; Verdonck, Marc Gustaaf Celine;
 Van Brandt, Sven Franciscus Anna; Angibaud, Patrick
 Rene; Meerpoel, Lieven; Dyatkin, Alexey Borisovich
 Janssen Pharmaceutica N.V., Belg.
 PATENT ASSIGNEE(S):
 SOURCE: PCT Int. Appl., 52 pp.
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 8
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003075929	A1	20030918	WO 2003-EP2515	20030311
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
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AU 2003218737	A1	20030922	AU 2003-218737	20030311
AU 2003218737	B2	20080410		
EP 1485099	A1	20041215	EP 2003-711981	20030311
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
BR 2003007624	A	20050111	BR 2003-7624	20030311
CN 1639125	A	20050713	CN 2003-805675	20030311
CN 1642551	A	20050720	CN 2003-805833	20030311
JP 20055523379	T	20050825	JP 2003-574203	20030311
NZ 534832	A	20050930	NZ 2003-534832	20030311
CN 101007803	A	20070801	CN 2007-10005212	20030311
IN 2004DN02537	A	20070112	IN 2004-DN2537	20040831
ZA 2004007237	A	20050928	ZA 2004-7237	20040909
ZA 2004007235	A	20051004	ZA 2004-7235	20040909
ZA 2004007232	A	20051006	ZA 2004-7232	20040909
ZA 2004007233	A	20051006	ZA 2004-7233	20040909
ZA 2004007234	A	20051006	ZA 2004-7234	20040909
ZA 2004007236	A	20051006	ZA 2004-7236	20040909
MX 2004PA08797	A	20041126	MX 2004-PA8797	20040910
US 20050096468	A1	20050505	US 2004-507785	20040913
NO 2004004113	A	20040928	NO 2004-4113	20040928
PRIORITY APPLN. INFO.:				
		US 2002-363799P	P	20020313
		WO 2002-EP14833	A	20021223
		CN 2003-805921	A3	20030311
		WO 2003-EP2515	W	20030311

OTHER SOURCE(S): MARPAT 139:261308



AB This invention comprises aryl and heteroaryl hydroxamic acids (shown as I; variables defined below; e.g. II) having histone deacetylase inhibiting enzymic activity; their preparation, compns. containing them and their use as a medicine. Compds. I show excellent in-vitro histone deacetylase inhibiting enzymic activity, have advantageous properties with regard to cellular activity and specific properties with regard to inhibition of cell cycle progression at both G1 and G2 checkpoints (p21 induction capacity), and show good metabolic stability and high bioavailability and more particular show oral bioavailability. They can also be used for detection and identification of histone deacetylase. General synthetic procedures and characterization data for twenty-seven I are included; also, preps. of 12 intermediates are included. For example, a 59 % yield of 2-(4-(dimethylaminosulfonyl)piperazin-1-yl)pyrimidine-5-carbohydroxamic acid was obtained by removing the O-tetrahydropyranyl group of its ester using trifluoroacetic acid; the ester was prepared in 61 % yield from N¹-(ethylcarboximidoyl)-N,N-dimethyl-3-propanediamine monohydrochloride, sodium 2-[4-(dimethylaminosulfonyl)piperazin-1-yl]pyrimidine-5-carboxylate, O-(tetrahydro-2H-pyran-2-yl)hydroxylamine, and 1-hydroxy-1H-benzotriazole in CH₂C₁₂/THF. The sodium salt was obtained by base hydrolysis of the Et ester; the ester was prepared in 73 % yield from Et 2-(piperazin-1-yl)pyrimidine-5-carboxylate and dimethylsulfamoyl chloride; Et 2-(piperazin-1-yl)pyrimidine-5-carboxylate was obtained in <96 % yield from Et 2-(4-benzylpiperazin-1-yl)pyrimidine-5-carboxylate by hydrogenation using Pd/C; the benzyl derivative was obtained from 1-(phenylmethyl)piperazine, (135 mL) was added gradually to a solution of potassium carbonate (0.18 mol) and 2-(methylsulfonyl)-5-pyrimidinecarboxylic acid Et ester, K₂CO₃ in MeCN. For I: n is 0-3; Q, X and Y are N or C; Z is N or CH; R¹ is -C(O)NR5R6, -N(H)C(O)R⁷, -C(O)-C1-6alkanediylR⁷, -NR8C(O)N(OH)R⁷, -NR8C(O)C1-6alkanediylR⁷, -NR8C(O)C:N(OH)R⁷ or another Zn-chelating-group; R² is H, halo, hydroxy, amino, nitro, C1-6alkyl, C1-6alkyloxy, trifluoromethyl, di(C1-6-alkyl)amino, hydroxymino or naphthalenylsulfonylpiperazinyl. R³ is H, C1-6-alkyl, arylC2-6alkanediyl, furanylcarbonyl, naphthalenylcarbonyl, -C(O)phenylR⁹, C1-6alkylaminocarbonyl, aminosulfonyl, arylaminosulfonyl, aminosulfonylamino, di(C1-6-alkyl)aminosulfonylamino, arylaminosulfonylamino, aminosulfonylaminoCl-6-alkyl, arylaminosulfonylaminoCl-6alkyl, di(C1-6-alkyl)aminocl-6alkyl, C11-12-alkylsulfonyl, di(C1-6-alkyl)aminosulfonyl, trihaloC1-6-alkylsulfonyl, di(aryl)C1-6alkylcarbonyl, thiophenylC1-6alkylcarbonyl, pyridinylcarbonyl or arylC1-6alkylcarbonyl. R⁴ is H, hydroxy, amino, hydroxyC1-6alkyl, Cl-6alkyl, Cl-6alkyloxy,

arylCl-6-alkyl, aminocarbonyl, hydroxycarbonyl, aminoCl-6-alkyl, aminocarbonylCl-6-alkyl, hydroxycarbonylCl-6-alkyl, hydroxyaminocarbonyl, Cl-6-alkyloxycarbonyl, Cl-6-alkylaminoCl-6-alkyl or di(Cl-6-alkyl)aminoCl-6-alkyl; when R3 and R4 are present on the same C atom, R3 and R4 together may form -C(O)-NH-CH₂-NR10- wherein R10 is H or aryl; when R3 and R4 are present on adjacent C atoms, R3 and R4 together may form :CH-CH:CH-CH: ; addnl. details are given in the claims.

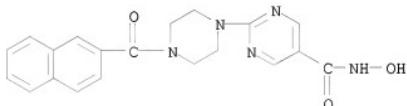
IT 603991-96-4P

RL: ARG (Analytical reagent use); PAC (Pharmacological activity); PKT (Pharmacokinetics); SPN (Synthetic preparation); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate and reagent for detection/identification of histone deacetylase; preparation of aryl and heteroaryl hydroxamic acids as inhibitors of histone deacetylase for treating proliferative diseases)

RN 603991-96-4 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-naphthalenylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)



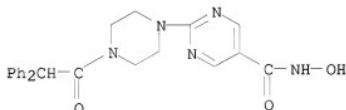
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603992-26-3P 603992-27-4P 603992-28-5P

RL: ARG (Analytical reagent use); PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate and reagent for detection/identification of histone deacetylase; preparation of aryl and heteroaryl hydroxamic acids as inhibitors of histone deacetylase for treating proliferative diseases)

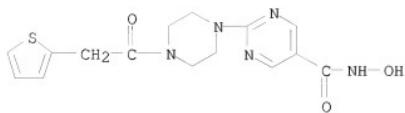
RN 603991-95-3 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-(2,2-diphenylacetyl)-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



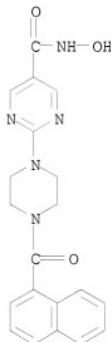
RN 603992-24-1 CAPLUS

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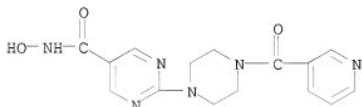
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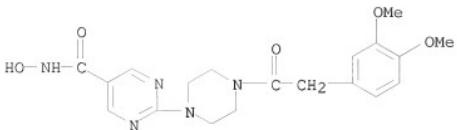
RN 603992-26-3 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(3-pyridinylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)



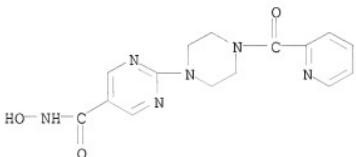
RN 603992-27-4 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[2-(3,4-dimethoxyphenyl)acetyl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



RN 603992-28-5 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-pyridinylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)

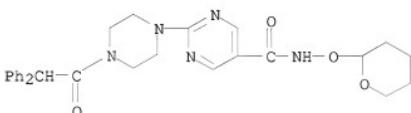


IT 603992-32-1P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (preparation of aryl and heteroaryl hydroxamic acids as inhibitors of histone deacetylase for treating proliferative diseases)

RN 603992-32-1 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-(2,2-diphenylacetyl)-1-piperazinyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]- (CA INDEX NAME)



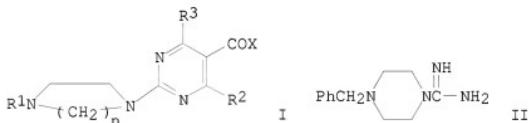
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THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 11 OF 11 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1986:442843 CAPLUS
 DOCUMENT NUMBER: 105:42843
 ORIGINAL REFERENCE NO.: 105:7101a,7104a
 TITLE: Pyrimidinylpiperazines
 INVENTOR(S): Kihara, Noriaki; Ishida, Tatsukazu; Isayama, Shigeru;
 Ishitoku, Takeshi; Tan, Hiroaki; Takahashi, Katsuya
 PATENT ASSIGNEE(S): Mitsui Petrochemical Industries, Ltd., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 28 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 61043173	A	19860301	JP 1984-163771	19840806
JP 05022702	B	19930330		
PRIORITY APPLN. INFO.:			JP 1984-163771	19840806
GI				



AB The title compds. [I, R1 = H, substituted Me, alkoxy carbonyl; R2, R3 = H, substituted alkyl; X = alkoxy, OH, (substituted) NH2; n = 2, 3], useful as herbicides against common weeds (no data), were prepared. Thus, the piperazinecarboxamide derivative II sulfate reacted with MeOCH:C(COME)CO2Me in MeOH/aqueous NaOH at room temperature overnight to give 88% I (R1 = PhCH2, n = 2,

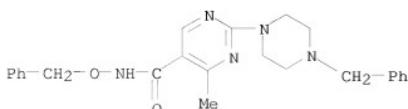
R2 = H, R3 = Me, X = OMe).

IT 102976-25-0P 102976-32-9P

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of, as herbicide)

RN 102976-25-0 CAPLUS

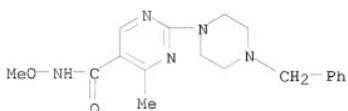
CN 5-Pyrimidinecarboxamide, 4-methyl-N-(phenylmethoxy)-2-[4-(phenylmethyl)-1-piperazinyl]- (CA INDEX NAME)



10/513699

RN 102976-32-9 CAPLUS

CN 5-Pyrimidinecarboxamide, N-methoxy-4-methyl-2-[4-(phenylmethyl)-1-piperazinyl]- (CA INDEX NAME)



10/513699

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COST IN U.S. DOLLARS                               SINCE FILE      TOTAL
                                                ENTRY SESSION
FULL ESTIMATED COST                           60.43    239.00

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)      SINCE FILE      TOTAL
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CA SUBSCRIBER PRICE                            -8.80    -8.80
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DICTIONARY FILE UPDATES: 14 SEP 2008 HIGHEST RN 1049627-95-31

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<http://www.cas.org/support/stnqgen/stndoc/properties.html>

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 ring nodes :
 5 6 7 8 9 10
 chain bonds :
 1-4 5-11
 ring bonds :
 5-6 5-7 6-8 7-9 8-10 9-10
 exact/norm bonds :
 1-4 5-6 5-7 5-11 6-8 7-9 8-10 9-10

G1:C,N

Match level :
 1:Atom 2:Atom 4:CLASS 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:CLASS
 Generic attributes :
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 Number of Carbon Atoms : less than 7
 Type of Ring System : Monocyclic

 Element Count :
 Node 1: Limited
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 N,N0-3

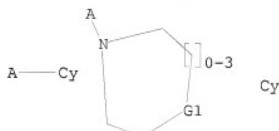
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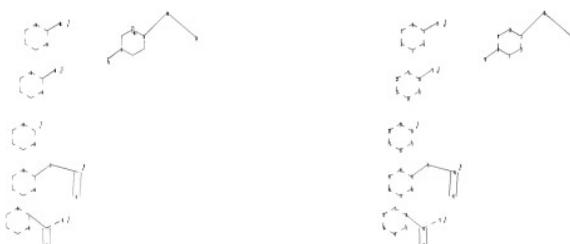


G1 C,N

Structure attributes must be viewed using STN Express query preparation.

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ring nodes :

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chain bonds :

10/513699

5-19 8-34 11-60 24-32 43-45 45-46 46-47 54-56 56-57 56-58 60-61
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54-56
isolated ring systems :
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G2:Ak,NH2,NO2

G3:O

G4: [*1], [*2], [*3], [*4], [*5]

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom
11:Atom 12:Atom 19:CLASS 20:Atom 21:Atom 22:Atom 23:Atom 24:Atom 25:Atom
26:Atom 27:Atom 28:Atom 29:Atom 30:Atom 31:Atom 32:CLASS 34:CLASS 39:Atom
40:Atom 41:Atom 42:Atom 43:Atom 44:Atom 45:CLASS 46:CLASS 47:CLASS 49:Atom
50:Atom 52:Atom 53:Atom 54:Atom 55:Atom 56:CLASS 57:CLASS 58:CLASS 60:CLASS
61:Atom

L5 STRUCTURE uploaded

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L5 HAS NO ANSWERS
L5 STR

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

Structure attributes must be viewed using STN Express query preparation.

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SEARCH TIME: 00.00.08

L6 10223 SEA SSS FUL L5

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COST IN U.S. DOLLARS SINCE FILE TOTAL

10/513699

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FULL ESTIMATED COST	179.28	418.28
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CA SUBSCRIBER PRICE	ENTRY	SESSION
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FILE COVERS 1907 - 15 Sep 2008 VOL 149 ISS 12
FILE LAST UPDATED: 14 Sep 2008 (20080914/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2008.

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

<http://www.cas.org/legal/infopolicy.html>

=> s 16 full
L7 4042 L6

=> s 17 and py<2003
22958911 PY<2003
L8 2880 L7 AND PY<2003

	SINCE FILE	TOTAL
COST IN U.S. DOLLARS	ENTRY	SESSION
FULL ESTIMATED COST	3.56	421.84
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
CA SUBSCRIBER PRICE	ENTRY	SESSION
	0.00	-8.80

FILE 'REGISTRY' ENTERED AT 16:06:14 ON 15 SEP 2008
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10/513699

STRUCTURE FILE UPDATES: 14 SEP 2008 HIGHEST RN 1049627-95-3
DICTIONARY FILE UPDATES: 14 SEP 2008 HIGHEST RN 1049627-95-3

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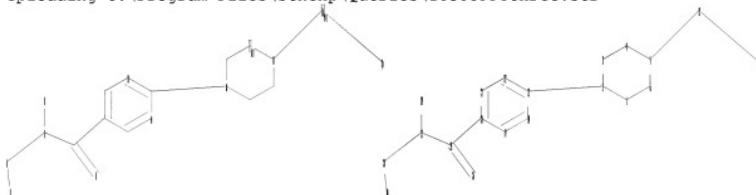
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<http://www.cas.org/support/stngen/stndoc/properties.html>

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ring nodes :
1 2 3 4 5 6 19 20 21 22 23 24
chain bonds :
2-23 5-13 13-14 20-25 25-26 25-27 27-28 27-29 28-30
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 19-20 19-24 20-21 21-22 22-23 23-24
exact/norm bonds :
1-2 1-6 2-3 2-23 3-4 4-5 5-6 5-13 13-14 25-26 25-27 27-28
exact bonds :
20-25 27-29 28-30
normalized bonds :
19-20 19-24 20-21 21-22 22-23 23-24
isolated ring systems :
containing 1 :

G1:C,N

G2:Ak,NH2,NO2

G3:O

10/513699

G4

G5:C,N,Zn,H

Match level :

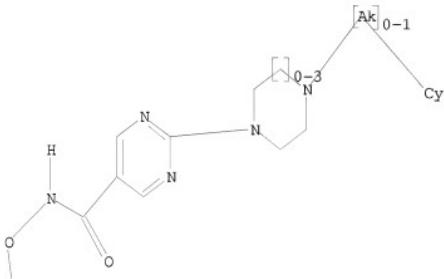
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21:Atom 22:Atom 23:Atom 24:Atom 25:CLASS 26:CLASS 27:CLASS 28:CLASS
29:CLASS 30:CLASS

L9 STRUCTURE UPLOADED

=> d 19

L9 HAS NO ANSWERS

L9 STR



G1 C,N

G2 Ak,NH2,NO2

G3 O

G4

G5 C,N,Zn,H

Structure attributes must be viewed using STN Express query preparation.

=> s 19 full

FULL SEARCH INITIATED 16:07:27 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 433 TO ITERATE

100.0% PROCESSED 433 ITERATIONS
SEARCH TIME: 00.00.01

112 ANSWERS

L10 112 SEA SSS FUL L9

=> file caplus

10/513699

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	178.82	600.66
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	0.00	-8.80

FILE 'CAPLUS' ENTERED AT 16:07:34 ON 15 SEP 2008
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FILE COVERS 1907 - 15 Sep 2008 VOL 149 ISS 12
FILE LAST UPDATED: 14 Sep 2008 (20080914/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2008.

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

<http://www.cas.org/legal/infopolicy.html>

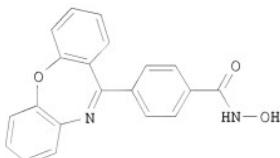
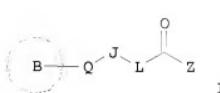
=> s l10 full
L11 13 L10

=> d ibib abs hitstr tot

L11 ANSWER 1 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2008:556979 CAPLUS
 DOCUMENT NUMBER: 148:538314
 TITLE: Preparation of tricyclic hydroxamic acids as
 inhibitors of histone deacetylase
 INVENTOR(S): Shapiro, Gideon; Moncuso, John; Pierre, Tessier; Leit,
 Silvana; Dexiel, Robert; David, Smil; Richard,
 Chesworth; Chantigny, Yves Andre; Patrick, Beaulieu
 PATENT ASSIGNEE(S): Methygene Inc., Can.; En Vivo Pharmaceuticals, Inc.
 SOURCE: PCT Int. Appl. 405pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008055068	A2	20080508	WO 2007-US82668	20071026
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
US 20080207590	A1	20080828	US 2007-925151	20071026
PRIORITY APPLN. INFO.:			US 2006-863347P	P 20061028
			US 2007-884287P	P 20070110

OTHER SOURCE(S): MARPAT 148:538314
 GI



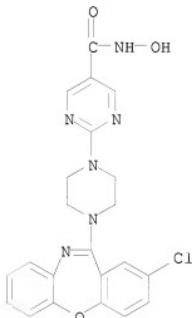
AB The title compds. I [Z = N(R1)OR2, H; L = a bond, N(OR2); when L = N(OR2), Z = H; when Z = H, L = N(OR2); R1, R2 = H, alkyl, aryl, etc.; J = a bond, :CH-, alkyl, alkyl(heteroalkyl)alkyl, etc.; Q = diazepine, pyrrolidine, diazabicyclo[3.3.1]nonane, etc.; B = dibenzob[f][1,4]oxazepine, benzo[b]pyrido[2,3-e][1,4]diazepine, benzo[f]thieno[2,3-b][1,4]oxazepine, etc.], useful for the inhibition of histone deacetylase, were prepared E.g., a 3-step synthesis of II, starting from 10,11-

dihydrodibenz[b,f][1,4]oxazepin-11-one, was given. All exemplified compds. I have an IC₅₀ of $\leq 10 \mu\text{M}$ against one or more of HDAC-1 through HDAC-11 (data for representative compds. I were given). Pharmaceutical composition comprising the compound I and methods of treating polyglutamine (polyQ) expansion diseases such as Huntington's disease, are disclosed.

IT 1024007-45-1P 1024009-50-4P 1024009-80-0P
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (preparation of tricyclic hydroxamic acids as inhibitors of histone deacetylase)

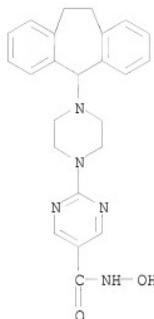
RN 1024007-45-1 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-(2-chlorodibenz[b,f][1,4]oxazepin-11-yl)-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



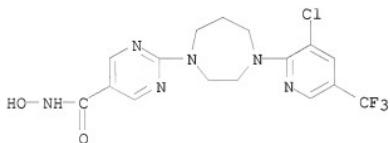
RN 1024009-50-4 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-(10,11-dihydro-5H-dibenzo[a,d]cyclohepten-5-yl)-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



RN 1024009-80-0 CAPLUS

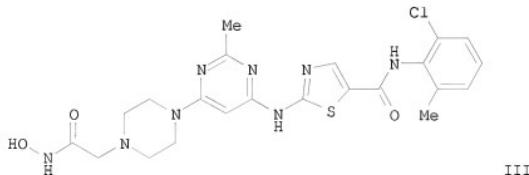
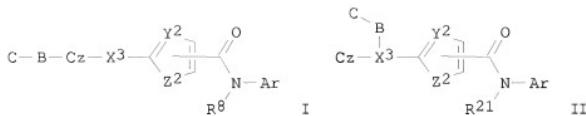
CN 5-Pyrimidinecarboxamide, 2-[4-(3-chloro-5-(trifluoromethyl)-2-pyridinyl)hexahydro-1H-1,4-diazepin-1-yl]-N-hydroxy- (CA INDEX NAME)



L11 ANSWER 2 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2008:353109 CAPLUS
 DOCUMENT NUMBER: 148:379651
 TITLE: Pyrimidine derivatives as tyrosine kinase inhibitors containing a zinc binding moiety and their preparation
 INVENTOR(S): Cai, Xiong; Qian, Changgeng; Gould, Stephen; Zhai, Haixiao
 PATENT ASSIGNEE(S): Curis, Inc., USA
 SOURCE: PCT Int. Appl., 81pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008033746	A2	20080320	WO 2007-US77970	20070910
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
US 20080125440	A1	20080529	US 2007-852450	20070910
PRIORITY APPLN. INFO.:			US 2006-843730P	P 20060911
			US 2007-895901P	P 20070320

OTHER SOURCE(S): MARPAT 148:379651
 GI



AB The invention relates to tyrosine kinase inhibitors of formula I and II that contain a zinc-binding moiety and their use in the treatment of tyrosine related diseases and disorders such as cancer. The said derivs. may further act as HDAC inhibitors. Compds. of formula I and II wherein Cz is (un)substituted (hetero)aryl, and (un)substituted heterocyclic; Ar is (un)substituted (hetero)aryl; X3 is NH, alkylamino, O, and S; Z2 is O, S, NH and alkylamino; Y2 is N, CH, C-halo, C-(hetero)aryl, etc.; R21 is H and aliphatic; B is a liner. C is urea, thiourea, acetyl, thioacetyl, etc.; R8 is H, acyl, and (un)substituted aliphatic group; and their geometric isomers, enantiomers, diastereoisomers, racemates, pharmaceutically acceptable salts, and solvates thereof, are claimed. Example compound III was prepared by a multistep procedure (procedure given). All the invention compds. were evaluated for their tyrosine kinase inhibitory activity.

IT 1012886-07-5P

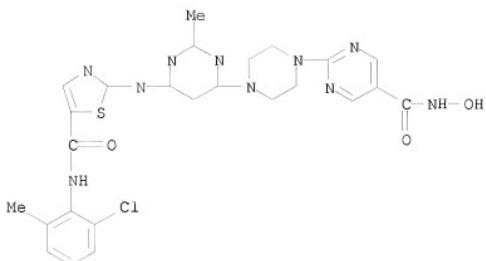
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of pyrimidine derivs. as tyrosine kinase inhibitors containing a zinc binding moiety)

RN 1012886-07-5 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[6-[[5-[(2-chloro-6-methylphenyl)amino]carbonyl]-2-thiazolyl]amino]-2-methyl-4-pyrimidinyl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

10/513699

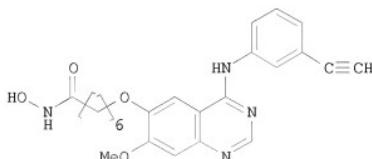


ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE

L11 ANSWER 3 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2008:353001 CAPLUS
DOCUMENT NUMBER: 148:355828
TITLE: Multi-functional small molecules as anti-proliferative
agents and their preparation
INVENTOR(S): Cai, Xiong; Qian, Changgeng; Gould, Stephen; Zhai,
Haixiao
PATENT ASSIGNEE(S): Curis, Inc., USA
SOURCE: PCT Int. Appl., 494pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008033747	A2	20080320	WO 2007-US77971	20070910
WO 2008033747	A9	20080724		
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RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GO, GW, ML, MR, NE, SN, TD, TG, TW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA			
US 20080221132	A1	20080911	US 2007-852458	20070910
PRIORITY APPLN. INFO.:			US 2006-843590P	P 20060911
			US 2007-895889P	P 20070320

OTHER SOURCE(S): MARPAT 148:355828
GI



A—B—C I

II

AB The invention relates to the compns., methods, and applications of an approach to selective inhibition of several cellular or mol. targets with a single small mol. More specifically, the present invention relates to multi-functional small mols. of formula I wherein one functionality is capable of inhibiting histone deacetylases (HDAC) and the other

functionality is capable of inhibiting a different cellular or mol. pathway involved in aberrant cell proliferation, differentiation or survival. Compds. of formula I wherein A is a pharmacophore of an anticancer agent capable of inhibiting at least one cellular or mol. pathway involved in the aberrant cell proliferation, differentiation or survival; B is a linker; C is a zinc-binding moiety; and their geometrical isomers, enantiomers, diastereoisomers, racemates, pharmaceutically acceptable salts, prodrugs and solvates thereof, are claimed. Example compound II was prepared by a multistep procedure (procedure given). All the invention compds. were evaluated for their antiproliferative activity (some data given).

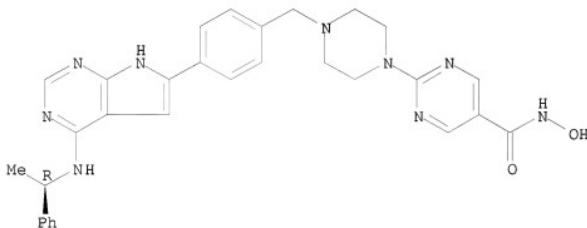
IT 1011716-90-7P 1012886-07-5P

RL: PAC (Pharmacological activity); PKT (Pharmacokinetics); PRPH (Prophetic); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (prophetic starting material; preparation of multi-functional small mols. as antiproliferative agents)

RN 1011716-90-7 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[[4-[4-[(1R)-1-phenylethyl]amino]-7H-pyrrolo[2,3-d]pyrimidin-6-yl]methyl]-1-piperazinyl]- (CA INDEX NAME)

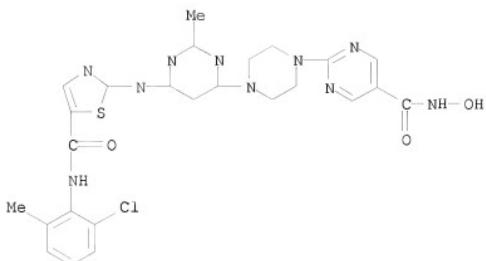
Absolute stereochemistry.



RN 1012886-07-5 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[6-[[5-[(2-chloro-6-methylphenyl)amino]carbonyl]-2-thiazolyl]amino]-2-methyl-4-pyrimidinyl]-1-piperazinyl-N-hydroxy- (CA INDEX NAME)

10/513699



ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE

L11 ANSWER 4 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2008:351928 CAPLUS
 DOCUMENT NUMBER: 148:355814
 TITLE: Preparation of (aralkylamino)(phenyl)pyrrolo[2,3-d]pyrimidine derivatives for use as protein tyrosine kinase (PTK) inhibitors
 INVENTOR(S): Cai, Xiong; Qian, Changgeng; Gould, Stephen
 PATENT ASSIGNEE(S): Curis, Inc., USA
 SOURCE: PCT Int. Appl., 123pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008033745	A2	20080320	WO 2007-US77968	20070910
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
US 20080161320	A1	20080703	US 2007-852440	20070910
PRIORITY APPLN. INFO.:			US 2006-843646P	P 20060911
			US 2007-895894P	P 20070320
OTHER SOURCE(S):	MARPAT	148:355814		
GI				

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB Fused bicyclic pyrimidine derivs. I and II [Ar = aryl, substituted arylheteroaryl or heteroaryl; Q = absent or (un)substituted alkyl; X = O, S, NH, or alkylamino; Z = O, S, NR1; Y = N or CR2; B = linker; D = C(O)NH2, NHC(S)CH3, CHC(O)NHacyl, etc.; R1 = H or (un)substituted alkyl; R2 = H, halo, (un)substituted aliphatic, aryl or heteroaryl], and their pharmaceutically acceptable salts, are prepared and disclosed as protein tyrosine kinase (PTK) inhibitors. Thus, e.g., III was prepared by N-alkylation of 1,4-dioxa-8-azaspiro[4.5]decane with 6-(4-(chloromethyl)phenyl)-N-((R)-1-phenylethyl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine (preparation given) and deprotection followed by condensation with 6-aminohexanoic acid Me ester and amidation with hydroxylamine. Select I were evaluated in EGFR assays, e.g., III demonstrated an IC50 value of ≤ 0.1 μ M.

IT 1011716-90-7P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES

10/513699

(Uses)

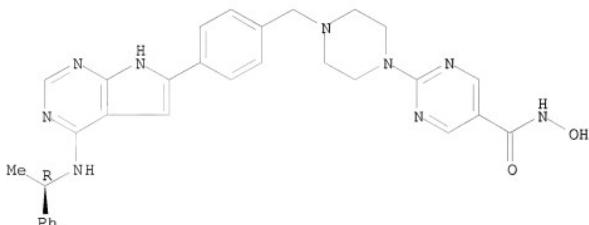
(preparation of (aralkylamino)(phenyl)pyrrolopyrimidine derivs. for use as protein tyrosine kinase (PTK) inhibitors)

RN 1011716-90-7 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[[4-[4-[[1R]-1-phenylethyl]amino]-

7H-pyrrolo[2,3-d]pyrimidin-6-ylphenyl)methyl]-1-piperazinyl)- (CA INDEX NAME)

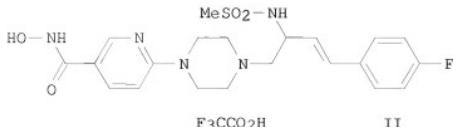
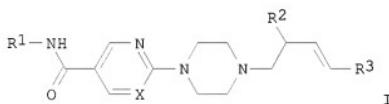
Absolute stereochemistry.



L11 ANSWER 5 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2007:816930 CAPLUS
 DOCUMENT NUMBER: 147:211903
 TITLE: Preparation of pyrimidine derivatives as histone deacetylase inhibitors
 INVENTOR(S): Marconnet-Decrane, Laurence Francoise Bernadette; Gaurrand, Sandrine Francoise Dominique; Angibaud, Patrick Rene
 PATENT ASSIGNEE(S): Janssen Pharmaceutica N.V., Belg.
 SOURCE: PCT Int. Appl., 62pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007082874	A1	20070726	WO 2007-EP50371	20070116
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JE, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, IJ, TM				
CA 2630717	A1	20070726	CA 2007-2630717	20070116
PRIORITY APPLN. INFO.:			EP 2006-100570	A 20060119
			WO 2007-EP50371	W 20070116

OTHER SOURCE(S): MARPAT 147:211903
 GI



AB The title compds. with general formula I [wherein R1 = OH or substituted phenyl; X = N or CH; R2 = amino, alkylamino, alkoxy, OH, etc.; R3 = (un)substituted Ph, naphthalene, or heterocycle] or N-oxide forms, pharmaceutically acceptable salts, or stereoisomeric forms thereof were prepared as histone deacetylase (HDAC) inhibitors for the treatment of proliferative diseases. For example, compound II was prepared in a multi-step synthesis. In vitro assay for inhibition of HDAC was performed to measure the inhibition of HDAC enzymic activity, and colorimetric assay was performed to determine cellular activity on A2780 tumor cells. II showed HDAC inhibitory and anti-proliferative activities in the above two assays with pIC50 values of 7.0 and 5.3, resp. Formulations containing I as active ingredients were also reported.

IT 944738-91-4P 944738-94-7P 944738-97-0P

944739-00-8P 944739-08-6P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of pyrimidine derivs. as histone deacetylase inhibitors)

RN 944738-91-4 CAPLUS

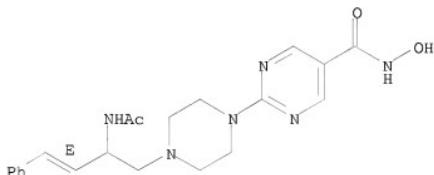
CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(acetylamino)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944738-90-3

CMF C21 H26 N6 O3

Double bond geometry as shown.



CM 2

CRN 76-05-1

CMF C2 H F3 O2



10/513699

RN 944738-94-7 CAPLUS

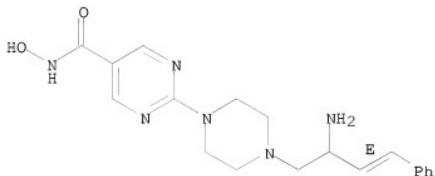
CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-amino-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944738-93-6

CMF C19 H24 N6 O2

Double bond geometry as shown.



CM 2

CRN 76-05-1

CMF C2 H F3 O2



RN 944738-97-0 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(2,5-dioxo-1-pyrrolidinyl)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

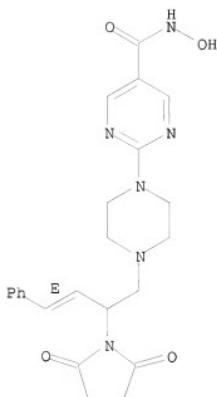
CM 1

CRN 944738-96-9

CMF C23 H26 N6 O4

Double bond geometry as shown.

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2



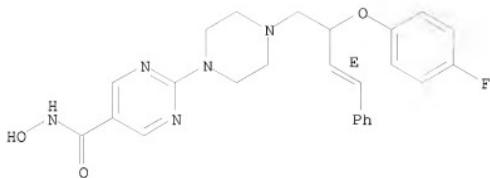
RN 944739-00-8 CAPLUS
CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(4-fluorophenoxy)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944738-99-2
CMF C25 H26 F N5 O3

Double bond geometry as shown.

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2

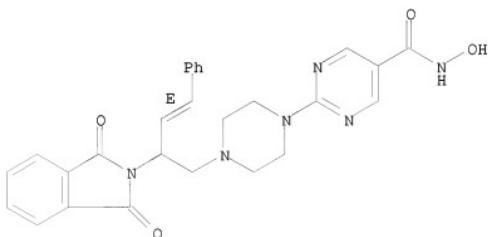


RN 944739-08-6 CAPLUS
CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(1,3-dihydro-1,3-dioxo-2H-isoindol-2-yl)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944739-07-5
CMF C27 H26 N6 O4

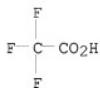
Double bond geometry as shown.



CM 2

10/513699

CRN 76-05-1
CMF C2 H F3 O2

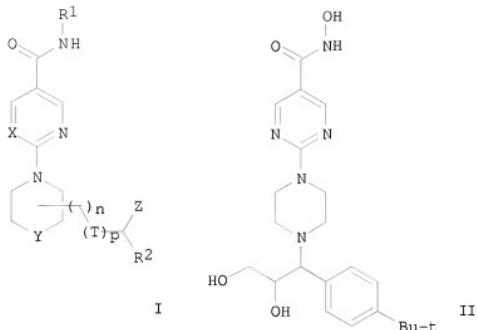


REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

10/513699

L11 ANSWER 6 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2007:816806 CAPLUS
DOCUMENT NUMBER: 147:211902
TITLE: Preparation of pyrimidine derivatives as histone deacetylase inhibitors
INVENTOR(S): Angibaud, Patrick Rene; Van Brandt, Sven Franciscus Anna; Marconnet-Decrane, Laurence Francoise Bernadette; Roux, Bruno
PATENT ASSIGNEE(S): Janssen Pharmaceutica N.V., Belg.
SOURCE: PCT Int. Appl., 63pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007082880	A1	20070726	WO 2007-EP50379	20070116
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JE, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, IJ, TM				
PRIORITY APPLN. INFO.: EP 2006-100571				A 20060119
OTHER SOURCE(S): MARPAT 147:211902				
GI				



AB The title compds. with general formula I (wherein R1 = OH or substituted phenyl; R2 = -CH2OH, -CH2COH3, -CH2COH2CH3, or -CH2CH(OH)CH2OH; T = N(R3), where R3 = H, alkyl, cycloalkyl, etc.; X = N or CH; Y = O, NH, CH2, etc.; n = 0-1; p = 0-1, provided that when p = 0 then n = 0 and Y = N, and -CH(R2)-Z is attached to Y; Z = (un)substituted aryl or heteroaryl) or N-oxide forms, pharmaceutically acceptable salts, or stereoisomeric forms thereof were prepared as histone deacetylase (HDAC) inhibitors for the treatment of proliferative diseases. For example, compound II was prepared in a multi-step synthesis. In vitro assay for inhibition of HDAC was performed to measure the inhibition of HDAC enzymic activity, and colorimetric assay was performed to determine cellular activity on A2780 tumor cells. II showed HDAC inhibitory and anti-proliferative activities in the above two assays with pIC50 values of 7.0 and 7.1, resp. Formulations containing I as active ingredients were also reported.

IT 944712-03-2P 944712-05-4P 944712-07-6P

944712-09-8P 944712-10-1P 944712-12-3P

~~944712-14=5P 944712-16=7P 944712-18=9P~~

RI: PAC (Pharmacological activity): SPM

RE: PAC (Pharmacological activity); SIR (Synthetic preparation); THER (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of pyrimidine derivs. as histone deacetylase inhibitors)

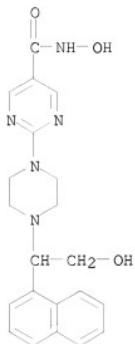
RN 944712-03-2 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-hydroxy-1-(1-naphthalenyl)ethyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-02-1
CMF C21 H23 N5 Q3

10/513699



CM 2

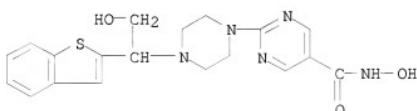
CRN 76-05-1
CMF C₂ H F₃ O₂



RN 944712-05-4 CAPLUS
CN 5-Pyrimidinecarboxamide, 2-[4-(1-benzo[b]thien-2-yl-2-hydroxyethyl)-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-04-3
CMF C₁₉ H₂₁ N₅ O₃ S



CM 2

10/513699

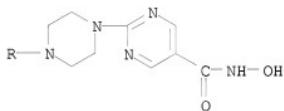
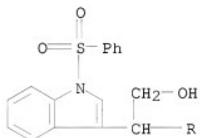
CRN 76-05-1
CMF C2 H F3 O2



RN 944712-07-6 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-hydroxy-1-[1-(phenylsulfonyl)-1H-indol-3-yl]ethyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-06-5
CMF C25 H26 N6 O5 S



CM 2

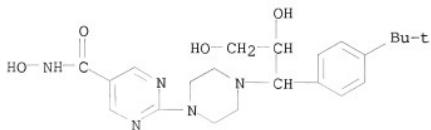
CRN 76-05-1
CMF C2 H F3 O2



RN 944712-09-8 CAPLUS
CN 5-Pyrimidinecarboxamide, 2-[4-[1-[4-(1,1-dimethylethyl)phenyl]-2,3-

10/513699

dihydroxypropyl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



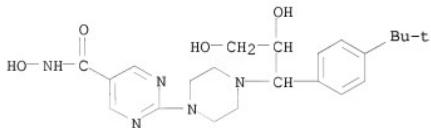
RN 944712-10-1 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1-[4-(1,1-dimethylethyl)phenyl]-2,3-dihydroxypropyl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1)
(CA INDEX NAME)

CM 1

CRN 944712-09-8

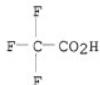
CMF C22 H31 N5 O4



CM 2

CRN 76-05-1

CMF C2 H F3 O2



RN 944712-12-3 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1(R,2S)-1-[4-(1,1-dimethylethyl)phenyl]-2,3-dihydroxypropyl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?)
(CA INDEX NAME)

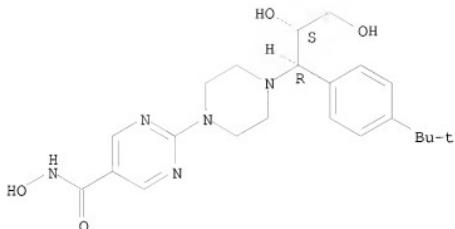
CM 1

CRN 944712-11-2

CMF C22 H31 N5 O4

Absolute stereochemistry.

10/513699



CM 2

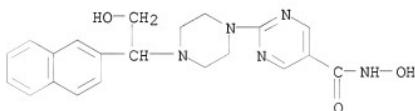
CRN 76-05-1
CMF C2 H F3 O2



RN 944712-14-5 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-hydroxy-1-(2-naphthalenyl)ethyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 944712-13-4
CMF C21 H23 N5 O3



CM 2

CRN 76-05-1
CMF C2 H F3 O2

10/513699

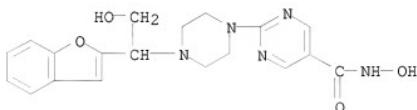


RN 944712-16-7 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1-(2-benzofuranyl)-2-hydroxyethyl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-15-6
CMF C19 H21 N5 O4



CM 2

CRN 76-05-1
CMF C2 H F3 O2



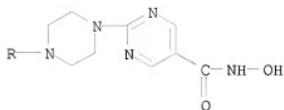
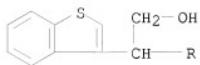
RN 944712-18-9 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-(1-benzo[b]thien-3-yl-2-hydroxyethyl)-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-17-8
CMF C19 H21 N5 O3 S

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2

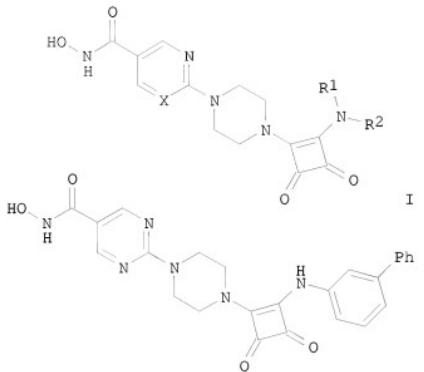


REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 7 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2007:485854 CAPLUS
 DOCUMENT NUMBER: 146:482095
 TITLE: Preparation of squaric acid derivatives as histone deacetylase (HDAC) inhibitors for the treatment of proliferative diseases
 INVENTOR(S): Van Emelen, Kristof
 PATENT ASSIGNEE(S): Janssen Pharmaceutica N. V., Belg.
 SOURCE: PCT Int. Appl., 37pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007048767	A1	20070503	WO 2006-EP67656	20061023
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, RU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
AU 2006307918	A1	20070503	AU 2006-307918	20061023
CA 2623360	A1	20070503	CA 2006-2623360	20061023
EP 1943232	A1	20080716	EP 2006-807466	20061023
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, RS				
PRIORITY APPLN. INFO.:			EP 2005-110080	A 20051027
			WO 2006-EP67656	W 20061023

OTHER SOURCE(S): MARPAT 146:482095
 GI



AB Title compds. I [wherein X = N or CH; R1, R2 = H, alkyl, Ph, etc.;] or N-oxides, pharmaceutically acceptable salts and stereoisomers thereof were prepared as histone deacetylase (HDAC) inhibitors. For instance, successive condensation of 3,4-diethoxy-3-cyclobutene-1,2-dione with 3-aminobiphenyl and 2-(1-piperazinyl)pyrimidine-5-carboxylic acid Et ester, ester hydrolysis, condensation of the resultant acid with NH₂O-THP, and deprotection with TFA gave hydroxamic acid II. This compds. showed inhibition against HDAC with pIC₅₀ = 7.7. The invented compds. are useful for the treatment of proliferative diseases.

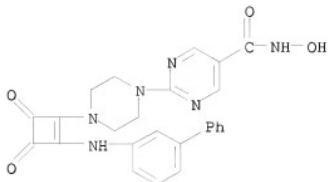
IT
935670-93-2P 935670-95-4P 935670-97-6P
935670-99-8P 935671-01-5P 935671-03-7P
935671-05-9P 935671-07-1P 935671-09-3P
935671-11-7P 935671-13-9P 935671-15-1P
935671-17-3P 935671-19-5P 935671-21-9P
935671-23-1P

RL: PAC (Pharmacological activity); **SPN:** (Synthetic preparation); **THU:** (Therapeutic use); **BIOL:** (Biological study); **PREP:** (Preparation); **USES** (Uses)

(preparation of squaric acid derivs. as histone deacetylase (HDAC) inhibitors for treatment of proliferative diseases)

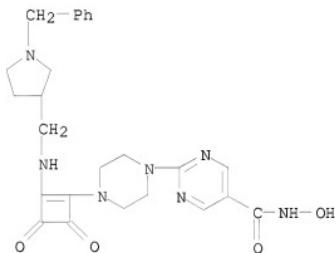
RN 935670-93-2 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[2-([1,1'-biphenyl]-3-ylamino)-3,4-dioxo-1-cyclobuten-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



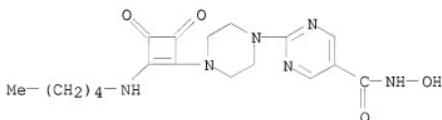
RN 935670-95-4 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3,4-dioxo-2-[(1-(phenylmethyl)-3-pyrrolidinyl]methyl]amino]-1-cyclobuten-1-yl]-1-piperazinyl-N-hydroxy- (CA INDEX NAME)



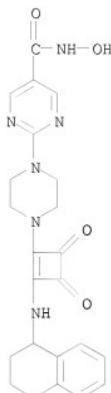
RN 935670-97-6 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3,4-dioxo-2-(pentylamino)-1-cyclobuten-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

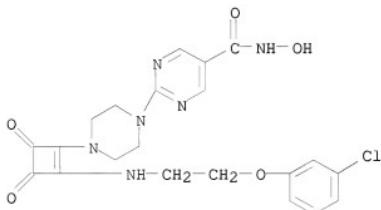


RN 935670-99-8 CAPLUS

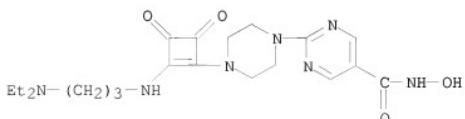
CN 5-Pyrimidinecarboxamide, 2-[4-[3,4-dioxo-2-[(1,2,3,4-tetrahydro-1-naphthalenyl)amino]-1-cyclobuten-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



RN 935671-01-5 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-{2-[(2-(3-chlorophenoxy)ethyl]amino]-3,4-dioxo-1-cyclobuten-1-yl}-N-hydroxy- (CA INDEX NAME)



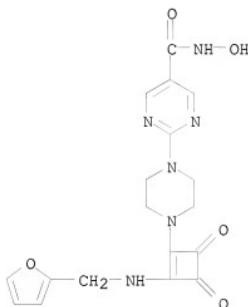
RN 935671-03-7 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-{2-[(3-(diethylamino)propyl]amino]-3,4-dioxo-1-cyclobuten-1-yl}-N-hydroxy- (CA INDEX NAME)



10/513699

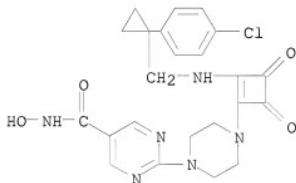
RN 935671-05-9 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[2-[(2-furanylmethyl)amino]-3,4-dioxo-1-cyclobuten-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



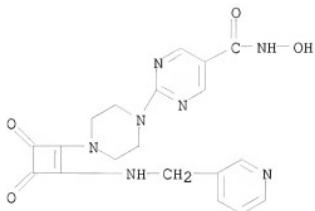
RN 935671-07-1 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[2-[[1-(4-chlorophenyl)cyclopropyl]methyl]amino]-3,4-dioxo-1-cyclobuten-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



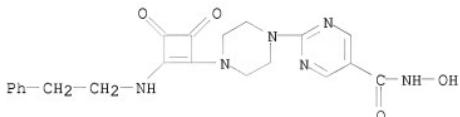
RN 935671-09-3 CAPLUS

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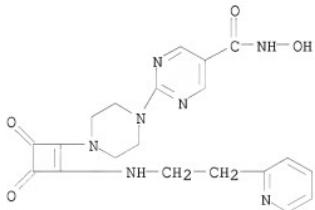
RN 935671-11-7 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3,4-dioxo-2-[(2-phenylethyl)amino]-1-cyclobuten-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



RN 935671-13-9 CAPLUS

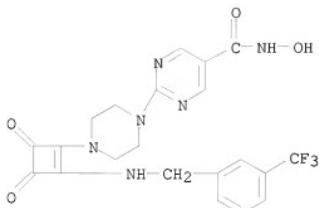
CN 5-Pyrimidinecarboxamide, 2-[4-[3,4-dioxo-2-[(2-(2-pyridinyl)ethyl)amino]-1-cyclobuten-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



RN 935671-15-1 CAPLUS

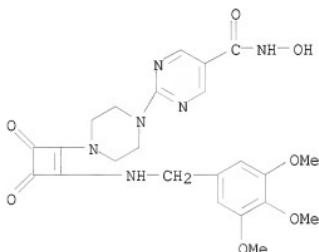
CN 5-Pyrimidinecarboxamide, 2-[4-[3,4-dioxo-2-[[[3-(trifluoromethyl)phenyl]methyl]amino]-1-cyclobuten-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

10/513699



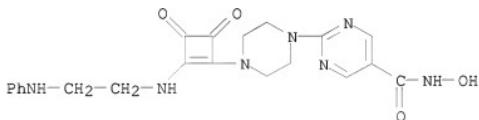
RN 935671-17-3 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3,4-dioxo-2-[(3,4,5-trimethoxyphenyl)methyl]amino]-1-cyclobuten-1-yl]-1-piperazinyl-N-hydroxy-
(CA INDEX NAME)



RN 935671-19-5 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3,4-dioxo-2-[(2-(phenylamino)ethyl]amino]-1-cyclobuten-1-yl]-1-piperazinyl-N-hydroxy-
(CA INDEX NAME)

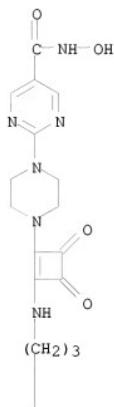


RN 935671-21-9 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3,4-dioxo-2-[(3-(2-oxo-1-pyrrolidinyl)propyl]amino]-1-cyclobuten-1-yl]-1-piperazinyl-N-hydroxy-
(CA INDEX NAME)

10/513699

PAGE 1-A

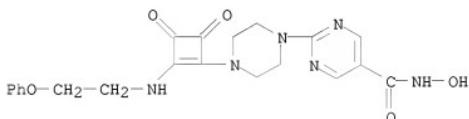


PAGE 2-A



RN 935671-23-1 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3,4-dioxo-2-[(2-phenoxyethyl)amino]-1-cyclobuten-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



REFERENCE COUNT:

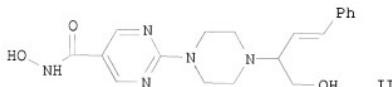
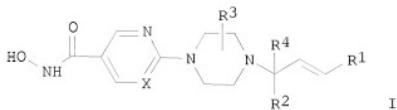
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THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 8 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2006:101446 CAPLUS
 DOCUMENT NUMBER: 144:192266
 TITLE: Preparation of substituted propenyl piperazine derivatives as novel inhibitors of histone deacetylase
 INVENTOR(S): Van Brandt, Sven Franciscus Anna; Van Emelen, Kristof;
 Angibaud, Patrick Rene; Marconnet-Decrane, Laurence
 Francoise Bernadette; Arts, Janine
 PATENT ASSIGNEE(S): Janssen Pharmaceutica N.V., Belg.
 SOURCE: PCT Int. Appl., 67 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006010749	A2	20060202	WO 2005-EP53611	20050725
WO 2006010749	A3	20060608		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
AU 2005266311	A1	20060202	AU 2005-266311	20050725
CA 2572971	A1	20060202	CA 2005-2572971	20050725
EP 1776358	A2	20070425	EP 2005-777776	20050725
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CN 1993356	A	20070704	CN 2005-80025487	20050725
JP 2008508234	T	20080321	JP 2007-523072	20050725
BR 2005013891	A	20080520	BR 2005-13891	20050725
KR 2007043978	A	20070426	KR 2007-701641	20070123
US 20070135424	A1	20070614	US 2007-626215	20070123
IN 2007DN00658	A	20070803	IN 2007-DN658	20070124
MX 200701119	A	20070315	MX 2007-1119	20070126
NO 200701117	A	20070227	NO 2007-1117	20070227
PRIORITY APPLN. INFO.:			EP 2004-77171	A 20040728
			US 2004-592357P	P 20040729
			WO 2005-EP53611	W 20050725

OTHER SOURCE(S): CASREACT 144:192266; MARPAT 144:192266
 GI



AB Substituted propenyl piperazine derivs. I, wherein X is independently N or CH; R1 is Ph, naphthalenyl or heterocyclyl; wherein each of said Ph or naphthalenyl is optionally substituted with one or two substituents each independently selected from halo, alkyl, alkyloxy, poly-halo-alkyl, aryl, hydroxy, cyano, amino, alkylcarbonylamino, alkylsulfonylamino, hydroxycarbonyl, alkyloxycarbonyl, hydroxyalkyl, alkyloxymethyl, aminomethyl, alkylaminomethyl, alkylcarbonylaminomethyl, alkylsulfonylaminomethyl, aminosulfonyl, alkylaminosulfonyl or heterocyclyl; R2 is hydrogen, -CH2R5, trifluoromethyl, -C(O)-R6, or -CH-NR/R8; wherein each R5 is independently hydrogen, hydroxy, alkyloxy, alkyloxyalkyloxy, alkylcarbonyloxy, piperazinyl, N-methylpiperazinyl, morpholinyl, thiomorpholinyl, imidazolyl or triazolyl; each R6 is independently hydroxy, alkyloxy, amino or mono- or di(alkyl)amino, cycloalkylamino, hydroxylalkylamino, piperazinyl, N-methylpiperazinyl, morpholinyl or thiomorpholinyl; each R7 and R8 are independently hydrogen, alkyl, alkylcarbonyl, alkylsulfonyl, or mono- or di(alkyl)aminosulfonyl; R3 is hydrogen, hydroxymethyl, aminomethyl or mono- or di(alkyl)aminomethyl; R4 is hydrogen or alkyl; were prepared and having histone deacetylase inhibiting enzymic activity and to inhibit proliferative conditions, such as cancer and psoriasis. Thus, propenyl piperazine derivative II was prepared and tested in vitro and in nude mice as inhibitor of histone deacetylase and was better than R306465 after oral administration. P21 enzyme linked immunosorbent assay has been applied to determine the p21 protein expression level in human A2780 ovarian carcinoma cells. In vitro assay for inhibition of histone deacetylase is reported. P21 induction was measured as the consequence of DNA damage or as the consequence of histone deacetylase inhibition. Antiproliferative activity of title compds. was determined on A2780 cells (neg. log value of the IC50, pIC50 = 7.9-8.2).

IT

875138-85-5P 875138-87-7P 875138-88-8P
 875138-89-9P 875138-90-2P 875138-91-3P
 875138-93-5P 875138-94-6P 875138-98-0P
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10/513699

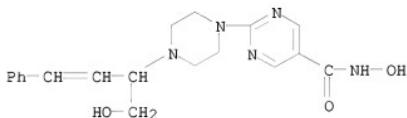
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RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of substituted propenyl piperazine derivs. as novel inhibitors of histone deacetylase)

RN 875138-85-5 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(1-(hydroxymethyl)-3-phenyl-2-propen-1-yl)-1-piperazinyl]- (CA INDEX NAME)



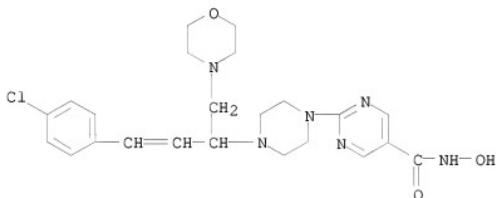
RN 875138-87-7 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3-(4-chlorophenyl)-1-(4-morpholinylmethyl)-2-propen-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-86-6

CMF C23 H29 Cl N6 O3



CM 2

CRN 76-05-1

CMF C2 H F3 O2

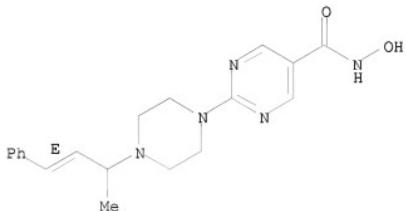


10/513699

RN 875138-88-8 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-methyl-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)

Double bond geometry as shown.



RN 875138-89-9 CAPLUS

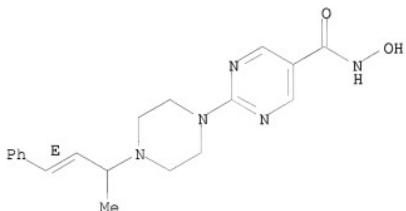
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-methyl-3-phenyl-2-propen-1-yl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-88-8

CMF C19 H23 N5 O2

Double bond geometry as shown.



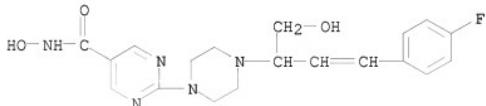
CM 2

CRN 76-05-1

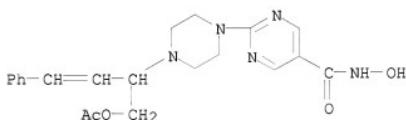
CMF C2 H F3 O2



RN 875138-90-2 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[3-(4-fluorophenyl)-1-(hydroxymethyl)-2-propen-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

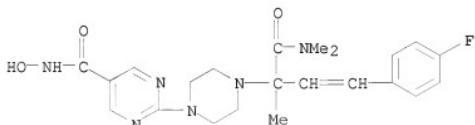


RN 875138-91-3 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(acetoxy)methyl]-3-phenyl-2-propen-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



RN 875138-93-5 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(dimethylamino)carbonyl]-3-(4-fluorophenyl)-1-methyl-2-propen-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-92-4
CMF C22 H27 F N6 O3

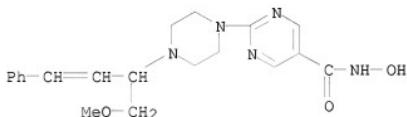
CM 2

10/513699

CRN 76-05-1
CMF C2 H F3 O2



RN 875138-94-6 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-(methoxymethyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)

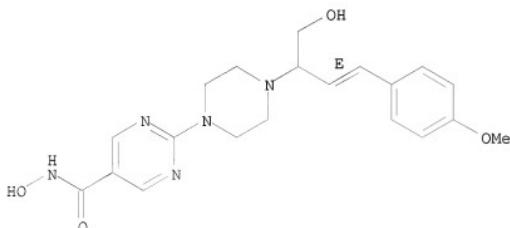


RN 875138-98-0 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-(hydroxymethyl)-3-(4-methoxyphenyl)-2-propen-1-yl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-97-9
CMF C20 H25 N5 O4

Double bond geometry as shown.



CM 2

CRN 76-05-1

10/513699

CMF C2 H F3 O2



RN 875139-00-7 CAPLUS

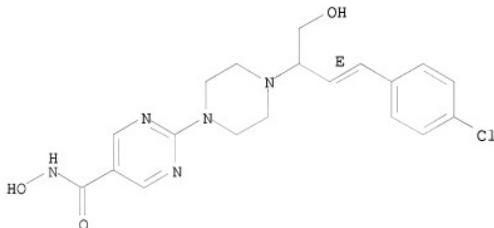
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CM 1

CRN 875138-99-1

CMF C19 H22 Cl N5 O3

Double bond geometry as shown.



CM 2

CRN 76-05-1

CMF C2 H F3 O2



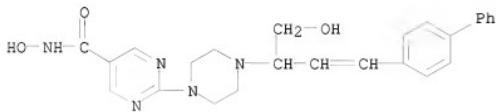
RN 875139-02-9 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3-[1,1'-biphenyl]-4-yl]-1-(hydroxymethyl)-2-propen-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

10/513699

CRN 875139-01-8
CMF C25 H27 N5 O3



CM 2

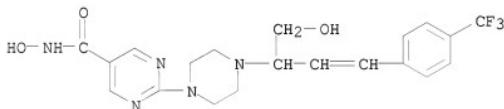
CRN 76-05-1
CMF C2 H F3 O2



RN 875139-04-1 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-(hydroxymethyl)-3-[4-(trifluoromethyl)phenyl]-2-propenyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-03-0
CMF C20 H22 F3 N5 O3



CM 2

CRN 76-05-1
CMF C2 H F3 O2

10/513699



RN 875139-06-3 CAPLUS

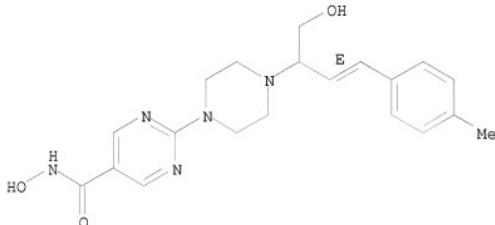
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-(hydroxymethyl)-3-(4-methylphenyl)-2-propen-1-yl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-05-2

CMF C20 H25 N5 O3

Double bond geometry as shown.



CM 2

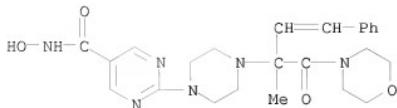
CRN 76-05-1

CMF C2 H F3 O2



RN 875139-07-4 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-methyl-1-(4-morpholinylcarbonyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)



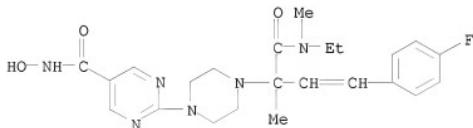
RN 875139-09-6 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(ethylmethylamino)carbonyl]-3-(4-fluorophenyl)-1-methyl-2-propenyl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-08-5

CMF C23 H29 F N6 O3



CM 2

CRN 76-05-1

CMF C2 H F3 O2



RN 875139-11-0 CAPLUS

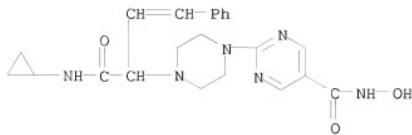
CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(cyclopropylamino)carbonyl]-3-phenyl-2-propenyl-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-10-9

CMF C22 H26 N6 O3

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2

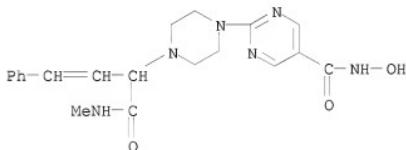


RN 875139-13-2 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(1-[(methylamino)carbonyl]-3-phenyl-2-propen-1-yl)-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-12-1
CMF C20 H24 N6 O3



CM 2

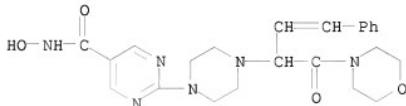
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10/513699



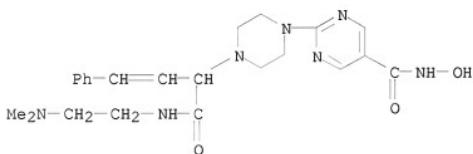
RN 875139-14-3 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-(4-morpholinylcarbonyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)



RN 875139-15-4 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(2-(dimethylamino)ethyl]amino]carbonyl]-3-phenyl-2-propen-1-yl]-1-piperazinyl-N-hydroxy- (CA INDEX NAME)



RN 875139-17-6 CAPLUS

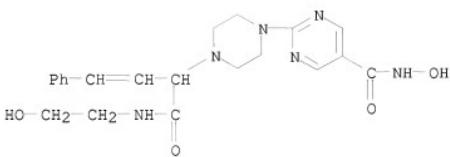
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-[(2-hydroxyethyl)amino]carbonyl]-3-phenyl-2-propen-1-yl]-1-piperazinyl-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-16-5

CMF C21 H26 N6 O4

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2

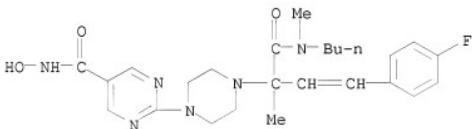


RN 875139-19-8 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[(butylmethylamino)carbonyl]-3-(4-fluorophenyl)-2-propenyl-1-yl]-1-piperazinyl-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-18-7
CMF C25 H33 F N6 O3

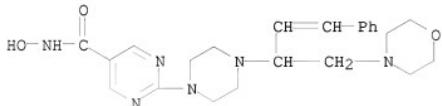


CM 2

CRN 76-05-1
CMF C2 H F3 O2

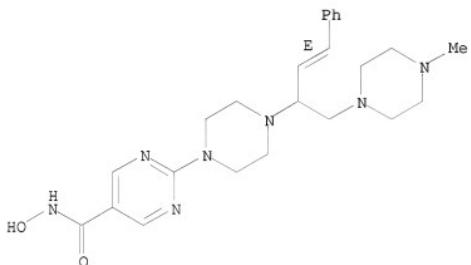


RN 875139-20-1 CAPLUS
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RN 875139-21-2 CAPLUS
 CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-[(4-methyl-1-piperazinyl)methyl]-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)

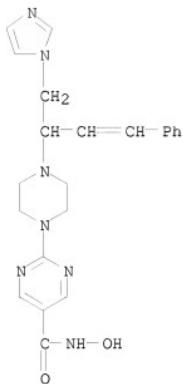
Double bond geometry as shown.



RN 875139-23-4 CAPLUS
 CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-(1H-imidazol-1-ylmethyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

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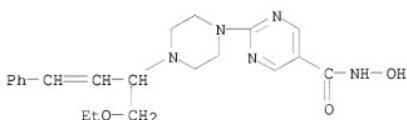
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CM 2

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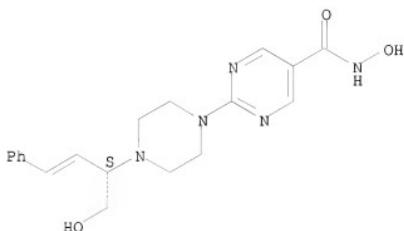


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Absolute stereochemistry.

10/513699

Double bond geometry unknown.

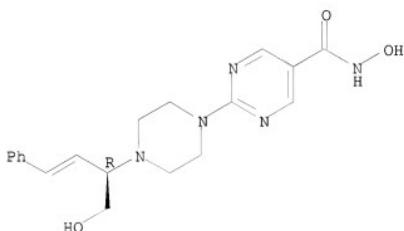


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Absolute stereochemistry.

Double bond geometry unknown.

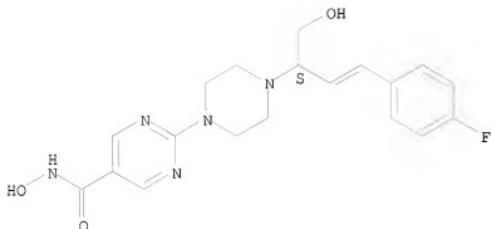


RN 875139-27-8 CAPLUS

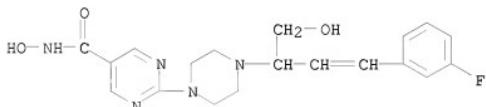
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Absolute stereochemistry.

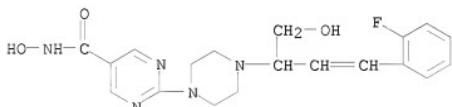
Double bond geometry unknown.



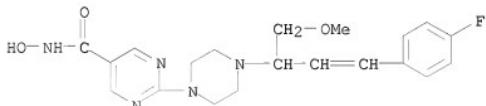
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RN 875139-29-0 CAPLUS
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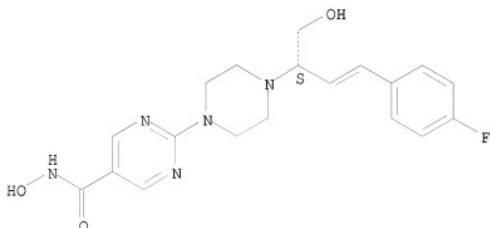


RN 875139-31-4 CAPLUS
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10/513699

propen-1-yl]-1-piperazinyl]-N-hydroxy-, hydrochloride (1:1) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.

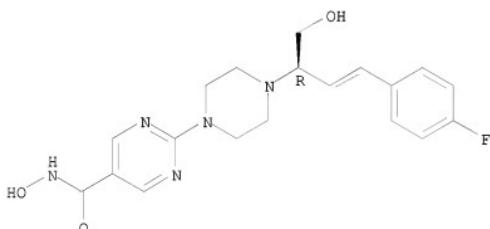


● HCl

RN 875139-69-8 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-((1R)-3-(4-fluorophenyl)-1-(hydroxymethyl)-2-propen-1-yl)-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.

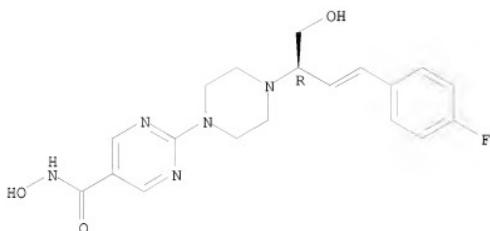


RN 875139-70-1 CAPLUS

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Absolute stereochemistry.
Double bond geometry unknown.

10/513699

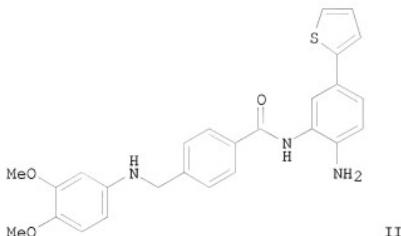
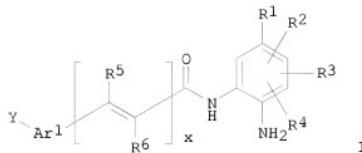


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L11 ANSWER 9 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2005:300395 CAPLUS
 DOCUMENT NUMBER: 142:355054
 TITLE: Preparation of amide derivatives as inhibitors of histone deacetylase
 INVENTOR(S): Moradei, Oscar; Paquin, Isabelle; Leit, Silvana;
 Frechette, Sylvie; Vaisburg, Arkadii; Besterman,
 Jeffrey M.; Tessier, Pierre; Mallais, Tammy C.
 PATENT ASSIGNEE(S): Methylgene, Inc., Can.
 SOURCE: PCT Int. Appl., 559 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005030705	A1	20050407	WO 2004-US31591	20040924
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			US 2004-561082P	P 20040409
			JP 2006-528279	A3 20040924
			WO 2004-US31591	W 20040924

OTHER SOURCE(S): CASREACT 142:355054; MARPAT 142:355054
 GI



AB Title compds. I [Arl = (un)saturated-, (un)substituted-mono or fused poly-cyclic hydrocarbyl optionally containing 1-4 heteroatoms per ring; R1 = (un)substituted-mono-, -bi-, -tri-cyclic-aryl or -heteroaryl; R2, R3, and R4 independently = H, halo, amino, etc.; R5 and R6 independently = H, alkyl, aryl, etc.; x = 0-1; Y = any pharmaceutically acceptable chemical moiety consisting of 1 to 50 atoms with provisions] and their pharmaceutically acceptable salts, are prepared and disclosed as inhibitors of histone deacetylase. Thus, e.g., II was prepared by Suzuki coupling of 2-bromo-2-nitro-phenylamine (preparation given) with 2-thiopheneboronic acid followed by carbonylation with 4-[3,4-dimethoxy-(phenylamino)-methyl]benzoic acid (preparation given) and subsequent reduction. The inhibitory capability of I towards antiproliferative activity of histone deacetylase enzyme was evaluated using 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT) assay and it revealed that certain compds. of the invention had MTT IC₅₀ values in the range of below 1 up to 20 μ M. I as histone deacetylase inhibitors should prove useful in the treatment of diseases such as, but not limited to, cell proliferative disease, protozoal disease, and fungal disease.

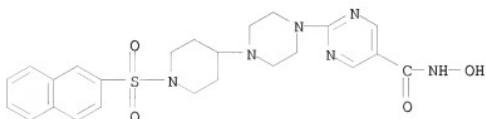
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RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

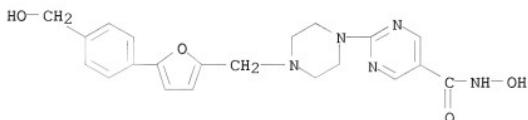
(preparation of amide derivs. as inhibitors of histone deacetylase)

10/513699

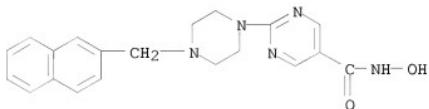
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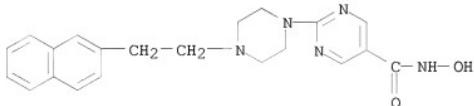
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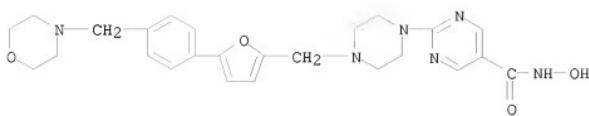
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RN 603985-94-0 CAPLUS
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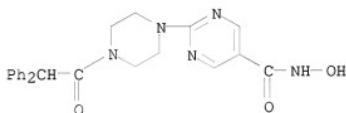
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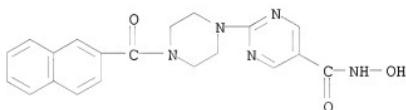
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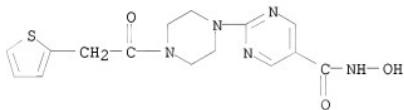
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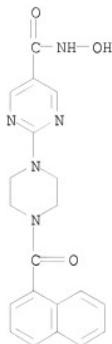
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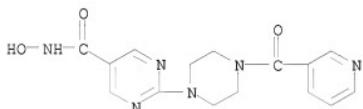


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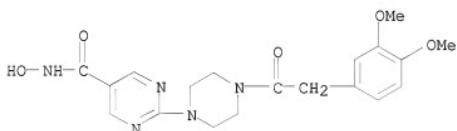
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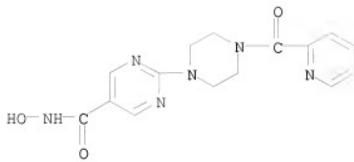


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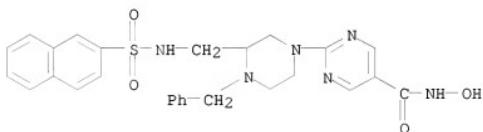
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10/513699



RN 604784-81-8 CAPLUS

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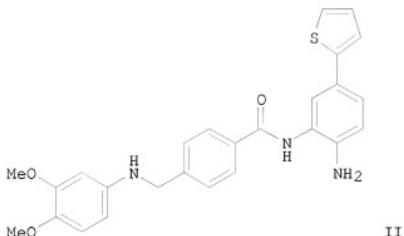
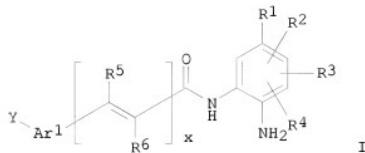
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THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 10 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2005:300394 CAPLUS
 DOCUMENT NUMBER: 142:373563
 TITLE: Preparation of amide derivatives as inhibitors of histone deacetylase
 INVENTOR(S): Moradei, Oscar; Paquin, Isabelle; Leit, Silvana;
 Frechette, Sylvie; Vaisburg, Arkadii; Besterman,
 Jeffrey M.; Tessier, Pierre; Mallais, Tammy C.
 PATENT ASSIGNEE(S): Methylgene, Inc., Can.
 SOURCE: PCT Int. Appl., 389 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005030704	A1	20050407	WO 2004-US31590	20040924
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RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, T2, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
JP 2008094847	A	20080424	JP 2007-281356 US 2003-505884P US 2003-532973P US 2004-561082P JP 2006-528279	20071030 P 20030924 P 20031229 P 20040409 A3 20040924
PRIORITY APPLN. INFO.:				
OTHER SOURCE(S): GI	CASREACT 142:373563; MARPAT 142:373563			



AB Title compds. I [Arl = (un)saturated-, (un)substituted-mono or fused poly-cyclic hydrocarbyl optionally containing 1-4 heteroatoms per ring; R1 = (un)substituted-mono-, -bi-, -tri-cyclic-aryl or -heteroaryl; R2, R3, and R4 independently = H, halo, amino, etc.; R5 and R6 independently = H, alkyl, aryl, etc.; x = 0-1; Y = any pharmaceutically acceptable chemical moiety consisting of 1 to 50 atoms with provisions] and their pharmaceutically acceptable salts, are prepared and disclosed as inhibitors of histone deacetylase. Thus, e.g., II was prepared by Suzuki coupling of 2-bromo-2-nitro-phenylamine (preparation given) with 2-thiopheneboronic acid followed by carbonylation with 4-[3,4-dimethoxy-(phenylamino)-methyl]benzoic acid (preparation given) and subsequent reduction. The inhibitory capability of I towards antiproliferative activity of histone deacetylase enzyme was evaluated using 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT) assay and it revealed that certain compds. of the invention had MTT IC₅₀ values in the range of below 1 up to 20 μ M. I as histone deacetylase inhibitors should prove useful in the treatment of diseases such as, but not limited to, cell proliferative disease, protozoal disease, and fungal disease.

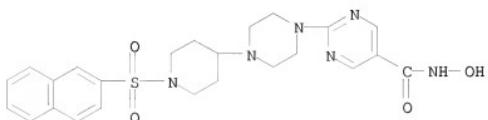
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RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

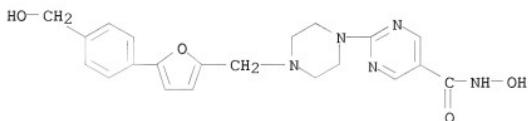
(preparation of amide derivs. as inhibitors of histone deacetylase)

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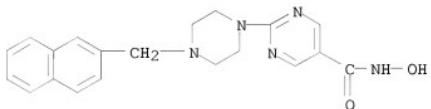
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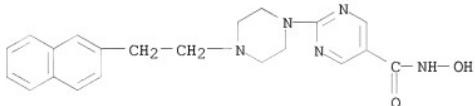
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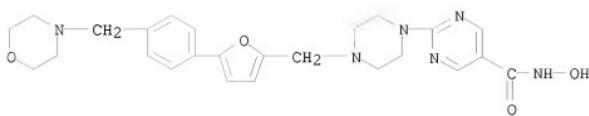
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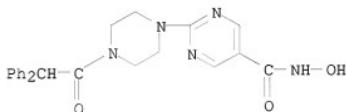
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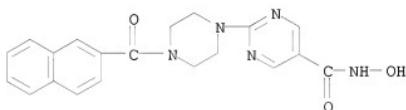
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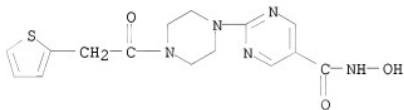
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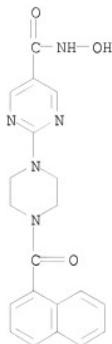
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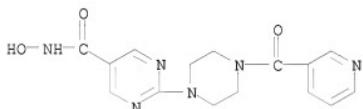


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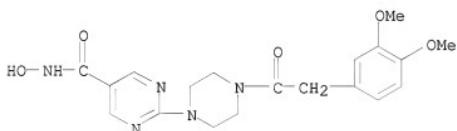
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CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(3-pyridinylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)

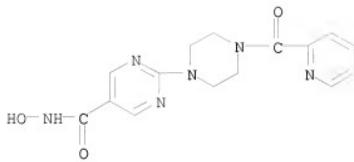


RN 603992-27-4 CAPLUS
CN 5-Pyrimidinecarboxamide, 2-[4-(2-(3,4-dimethoxyphenyl)acetyl)-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



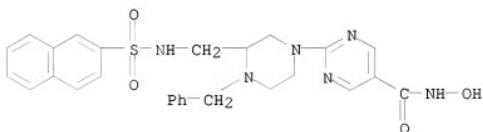
RN 603992-28-5 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-pyridinylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)

10/513699



RN 604784-81-8 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[3-[(2-naphthalenylsulfonyl)amino]methoxy]-4-(phenylmethyl)-1-piperazinyl- (CA INDEX NAME)



REFERENCE COUNT:

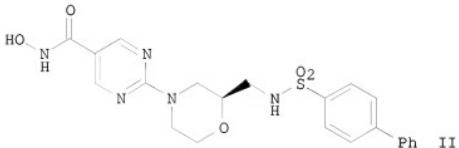
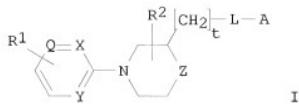
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THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 11 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2003:737757 CAPLUS
 DOCUMENT NUMBER: 139:276911
 TITLE: Preparation of N-(piperazinylmethyl-, piperidinylmethyl- and morpholinylmethyl) sulfonamides and amides as novel inhibitors of histone deacetylase
 INVENTOR(S): Van Emelen, Kristof
 PATENT ASSIGNEE(S): Janssen Pharmaceutica N.V., Belg.
 SOURCE: PCT Int. Appl., 69 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 8
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003076438	A1	20030918	WO 2003-EP2510	20030311
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KE, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2475766	A1	20030918	CA 2003-2475766	20030311
AU 2003218735	A1	20030922	AU 2003-218735	20030311
EP 1485378	A1	20041215	EP 2003-711979	20030311
EP 1485378	B1	20080618		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
BR 2003007606	A	20041221	BR 2003-7606	20030311
CN 1642948	A	20050720	CN 2003-805921	20030311
JP 2005526766	T	20050908	JP 2003-574655	20030311
NZ 534833	A	20060728	NZ 2003-534833	20030311
CN 101007803	A	20070801	CN 2007-10005212	20030311
AT 398615	T	20080715	AT 2003-711979	20030311
TW 283676	B	20070711	TW 2003-92105285	20030312
IN 2004DN02536	A	20070413	IN 2004-DN2536	20040831
US 20050165016	A1	20050728	US 2004-507084	20040908
MX 2004PA08795	A	20041126	MX 2004-PA8795	20040910
NO 2004004135	A	20040929	NO 2004-4135	20040929
PRIORITY APPLN. INFO.:			US 2002-363799P	P 20020313
			WO 2002-EP14833	A 20021223
			CN 2003-805921	A3 20030311
			WO 2003-EP2510	W 20030311

OTHER SOURCE(S): MARPAT 139:276911
 GI

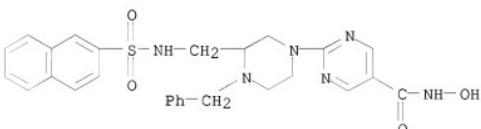


AB The title compds. [I; t = 0-4; Q, X, Y = N, C; Z = NH, O, CH₂; R1 = CONR₃R₄, NHCOR₇, CO(alkanediyl)SR₇, etc. (wherein R₃, R₄ = H, OH, alkyl, etc.; R₇ = H, alkyl, alkylcarbonyl, etc.); R2 = H, OH, NH₂, etc.; L = NR₉CO, NR₉SO₂, NR₉CH₂ (R₉ = H, alkyl, cycloalkyl, etc.); A = (un)substituted Ph, cycloalkyl, pyridyl, etc.] , having histone deacetylase inhibiting enzymic activity, were prepared and formulated. E.g., a multi-step synthesis of (+)-II which showed pIC₅₀ of 7.723 against HDAC, was given.

IT 604784-81-8P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of N-(piperazinylmethyl-, piperidinylmethyl- and morpholinylmethyl) sulfonamides and amides as novel inhibitors of histone deacetylase)
RN 604784-81-8 CAPLUS
CN 5-Pyridinylcarboxamide, N-hydroxy-2-[3-[(2-naphthalenylsulfonyl)amino]methoxy]-4-(phenylmethyl)-1-piperazinyl- (CA INDEX NAME)



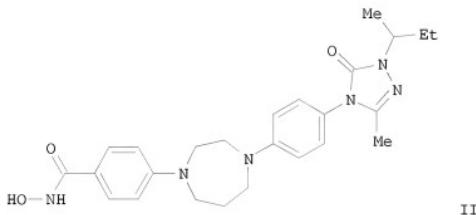
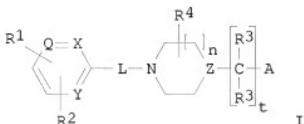
REFERENCE COUNT:

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THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE REFORMAT.

L11 ANSWER 12 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2003:737723 CAPLUS
 DOCUMENT NUMBER: 139:261309
 TITLE: Preparation of N-hydroxy-5-piperazino(piperidino or diazepino)-2-pyrimidinecarboxamides and N-hydroxy-4-piperazino(piperidino or diazepino)benzamides as new inhibitors of histone deacetylase
 INVENTOR(S): Angibaud, Patrick Rene; Pilatte, Isabelle Noelle Constance; Van Brandt, Sven Franciscus Anna; Roux, Bruno; Ten Holte, Peter; Verdonck, Marc Gustaaf Celine; Meerpoel, Lieven; Dyatkin, Alexey Borisovich Janssen Pharmaceutica N.V., Belg.
 PATENT ASSIGNEE(S):
 SOURCE: PCT Int. Appl., 72 pp.
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 8
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003076400	A1	20030918	WO 2003-EP2514	20030311
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2475764	A1	20030918	CA 2003-2475764	20030311
AU 2003218736	A1	20030922	AU 2003-218736	20030311
EP 1485353	A1	20041215	EP 2003-711980	20030311
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
BR 2003008081	A	20041221	BR 2003-8081	20030311
CN 1639125	A	20050713	CN 2003-805675	20030311
CN 1642551	A	20050720	CN 2003-805833	20030311
NZ 534834	A	20050729	NZ 2003-534834	20030311
JP 2005526067	T	20050902	JP 2003-574621	20030311
CN 101007803	A	20070801	CN 2007-10005212	20030311
IN 2004DN02533	A	20070413	IN 2004-DN2533	20040831
US 20050107384	A1	20050519	US 2004-506998	20040908
ZA 2004007237	A	20050928	ZA 2004-7237	20040909
ZA 2004007235	A	20051004	ZA 2004-7235	20040909
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ZA 2004007233	A	20051006	ZA 2004-7233	20040909
ZA 2004007234	A	20051006	ZA 2004-7234	20040909
ZA 2004007236	A	20051006	ZA 2004-7236	20040909
MX 2004PA08806	A	20041126	MX 2004-PA8806	20040910
NO 2004004194	A	20041001	NO 2004-4194	20041001
PRIORITY APPLN. INFO.:				
		US 2002-363799P	P	20020313
		WO 2002-EP14833	A	20021223
		CN 2003-805921	A3	20030311



AB The title compds. [I; n = 0-3; t = 0-4; Q, X, Y = N, C; Z = N, CH; R1 = CONR7R8, NHCOR9, CO(alkanediyl)SR9, etc. (wherein R7, R8 = H, OH, alkyl, etc.; R9 = H, alkyl, alkylcarbonyl, etc.); R2 = H, halo, OH, etc.; L = a bond, alkanediyl, alkanediyoxy, NH, CO, NHCO; each R3 = H and one H atom can be replaced by aryl; R4 = H, OH, NH2, etc.; A = (un)substituted Ph, cyclohexyl, pyridyl, etc.], having histone deacetylase inhibiting enzymic activity, were prepared and formulated. E.g., a multi-step synthesis of II which showed pIC50 of 5.121 against HDAC, was given.

IT 603985-83-7P 603985-87-1P 603985-89-3P

603985-91-7P 603985-95-1P

RL: PAC (Pharmacological activity); **SPN:** (Synthetic preparation); **THU:** (Therapeutic use); **BIOL:** (Biological study); **PREP:** (Preparation); **USES** (Uses)

(preparation of piperazino(piperidino or diazepino) substituted 2-pyrimidinecarboxylic acids and N-hydroxybenzamides as new inhibitors of histone deacetylase)

RN 603985-83-7 CAPLUS

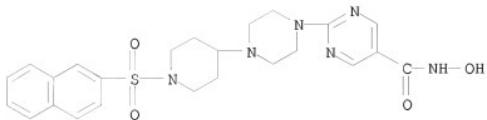
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-(2-naphthalenylsulfonyl)-4-piperidinyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (10:9) (CA INDEX NAME)

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CRN 603985-82-6

CMF C24 H28 N6 O4 S

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2

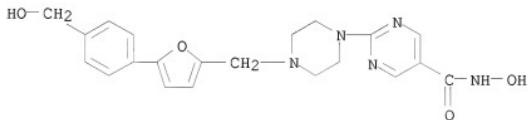


RN 603985-87-1 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[[5-[(4-hydroxymethyl)phenyl]-2-furanyl]methyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (5:4) (CA INDEX NAME)

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CRN 603985-86-0
CMF C21 H23 N5 O4



CM 2

CRN 76-05-1
CMF C2 H F3 O2



10/513699

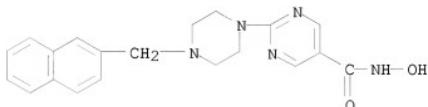
RN 603985-89-3 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-naphthalenylmethyl)-1-piperazinyl]-, 2,2,2-trifluoroacetate (5:4) (CA INDEX NAME)

CM 1

CRN 603985-88-2

CMF C20 H21 N5 O2



CM 2

CRN 76-05-1

CMF C2 H F3 O2



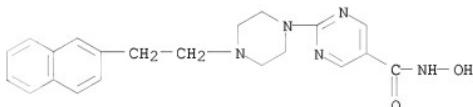
RN 603985-91-7 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-(2-naphthalenyl)ethyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (5:4) (CA INDEX NAME)

CM 1

CRN 603985-90-6

CMF C21 H23 N5 O2



CM 2

CRN 76-05-1

CMF C2 H F3 O2

10/513699



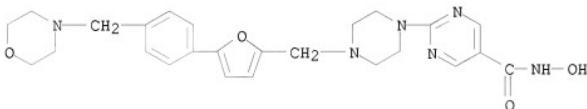
RN 603985-95-1 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(5-[4-(4-morpholinylmethyl)phenyl]-2-furanyl)methyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:2) (CA INDEX NAME)

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CRN 603985-94-0

CMF C25 H30 N6 O4



CM 2

CRN 76-05-1

CMF C2 H F3 O2



REFERENCE COUNT:

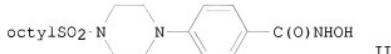
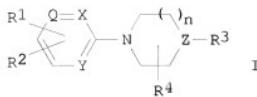
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THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 13 OF 13 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2003:737586 CAPLUS
 DOCUMENT NUMBER: 139:261308
 TITLE: Preparation of aryl and heteroaryl hydroxamic acids as inhibitors of histone deacetylase for treating proliferative diseases
 INVENTOR(S): Van Emelen, Kristof; Verdonck, Marc Gustaaf Celine;
 Van Brandt, Sven Franciscus Anna; Angibaud, Patrick
 Rene; Meerpoel, Lieven; Dyatkin, Alexey Borisovich
 Janssen Pharmaceutica N.V., Belg.
 PATENT ASSIGNEE(S):
 SOURCE: PCT Int. Appl., 52 pp.
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 8
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003075929	A1	20030918	WO 2003-EP2515	20030311
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2476065	A1	20030918	CA 2003-2476065	20030311
AU 2003218737	A1	20030922	AU 2003-218737	20030311
AU 2003218737	B2	20080410		
EP 1485099	A1	20041215	EP 2003-711981	20030311
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
BR 2003007624	A	20050111	BR 2003-7624	20030311
CN 1639125	A	20050713	CN 2003-805675	20030311
CN 1642551	A	20050720	CN 2003-805833	20030311
JP 20055523379	T	20050825	JP 2003-574203	20030311
NZ 534832	A	20050930	NZ 2003-534832	20030311
CN 101007803	A	20070801	CN 2007-10005212	20030311
IN 2004DN02537	A	20070112	IN 2004-DN2537	20040831
ZA 2004007237	A	20050928	ZA 2004-7237	20040909
ZA 2004007235	A	20051004	ZA 2004-7235	20040909
ZA 2004007232	A	20051006	ZA 2004-7232	20040909
ZA 2004007233	A	20051006	ZA 2004-7233	20040909
ZA 2004007234	A	20051006	ZA 2004-7234	20040909
ZA 2004007236	A	20051006	ZA 2004-7236	20040909
MX 2004PA08797	A	20041126	MX 2004-PA8797	20040910
US 20050096468	A1	20050505	US 2004-507785	20040913
NO 2004004113	A	20040928	NO 2004-4113	20040928
PRIORITY APPLN. INFO.:				
		US 2002-363799P	P	20020313
		WO 2002-EP14833	A	20021223
		CN 2003-805921	A3	20030311
		WO 2003-EP2515	W	20030311

OTHER SOURCE(S): MARPAT 139:261308



AB This invention comprises aryl and heteroaryl hydroxamic acids (shown as I; variables defined below; e.g. II) having histone deacetylase inhibiting enzymic activity; their preparation, compns. containing them and their use as a medicine. Compds. I show excellent in-vitro histone deacetylase inhibiting enzymic activity, have advantageous properties with regard to cellular activity and specific properties with regard to inhibition of cell cycle progression at both G1 and G2 checkpoints (p21 induction capacity), and show good metabolic stability and high bioavailability and more particular show oral bioavailability. They can also be used for detection and identification of histone deacetylase. General synthetic procedures and characterization data for twenty-seven I are included; also, preps. of 12 intermediates are included. For example, a 59 % yield of 2-(4-(dimethylaminosulfonyl)piperazin-1-yl)pyrimidine-5-carbohydroxamic acid was obtained by removing the O-tetrahydropyranyl group of its ester using trifluoroacetic acid; the ester was prepared in 61 % yield from N¹-(ethylcarboximidoyl)-N,N-dimethyl-3-propanediamine monohydrochloride, sodium 2-[4-(dimethylaminosulfonyl)piperazin-1-yl]pyrimidine-5-carboxylate, O-(tetrahydro-2H-pyran-2-yl)hydroxylamine, and 1-hydroxy-1H-benzotriazole in CH₂C₁₂/THF. The sodium salt was obtained by base hydrolysis of the Et ester; the ester was prepared in 73 % yield from Et 2-(piperazin-1-yl)pyrimidine-5-carboxylate and dimethylsulfamoyl chloride; Et 2-(piperazin-1-yl)pyrimidine-5-carboxylate was obtained in <96 % yield from Et 2-(4-benzylpiperazin-1-yl)pyrimidine-5-carboxylate by hydrogenation using Pd/C; the benzyl derivative was obtained from 1-(phenylmethyl)piperazine, (135 mL) was added gradually to a solution of potassium carbonate (0.18 mol) and 2-(methylsulfonyl)-5-pyrimidinecarboxylic acid Et ester, K₂CO₃ in MeCN. For I: n is 0-3; Q, X and Y are N or C; Z is N or CH; R1 is -C(O)NR5R6, -N(H)C(O)R7, -C(O)-C1-6alkanediylR7, -NR8C(O)N(OH)R7, -NR8C(O)C1-6alkanediylR7, -NR8C(O)C:N(OH)R7 or another Zn-chelating-group; R2 is H, halo, hydroxy, amino, nitro, Cl-6alkyl, Cl-6alkyloxy, trifluoromethyl, di(Cl-6-alkyl)amino, hydroxymino or naphthalenylsulfonylpiperazinyl. R3 is H, Cl-6-alkyl, arylC2-6alkanediyl, furanylcarbonyl, naphthalenylcarbonyl, -C(O)phenylR9, Cl-6alkylaminocarbonyl, aminosulfonyl, arylaminosulfonyl, aminosulfonylamino, di(Cl-6-alkyl)aminosulfonylamino, arylaminosulfonylamino, aminosulfonylaminoCl-6-alkyl, arylaminosulfonylaminoCl-6alkyl, di(Cl-6-alkyl)aminocl-6alkyl, Cl-12-alkylsulfonyl, di(Cl-6-alkyl)aminosulfonyl, trihaloCl-6-alkylsulfonyl, di(aryl)Cl-6alkylcarbonyl, thiophenylCl-6alkylcarbonyl, pyridinylcarbonyl or arylCl-6alkylcarbonyl. R4 is H, hydroxy, amino, hydroxyCl-6alkyl, Cl-6alkyl, Cl-6alkyloxy,

arylCl-6-alkyl, aminocarbonyl, hydroxycarbonyl, aminoCl-6-alkyl, aminocarbonylCl-6-alkyl, hydroxycarbonylCl-6-alkyl, hydroxyaminocarbonyl, Cl-6-alkyloxycarbonyl, Cl-6-alkylaminoCl-6-alkyl or di(Cl-6-alkyl)aminoCl-6-alkyl; when R3 and R4 are present on the same C atom, R3 and R4 together may form -C(O)-NH-CH₂-NR10- wherein R10 is H or aryl; when R3 and R4 are present on adjacent C atoms, R3 and R4 together may form :CH-CH:CH-CH: ; addnl. details are given in the claims.

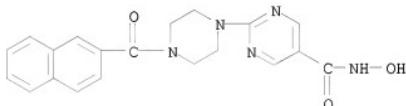
IT 603991-96-4P

RL: ARG (Analytical reagent use); PAC (Pharmacological activity); PKT (Pharmacokinetics); SPN (Synthetic preparation); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate and reagent for detection/identification of histone deacetylase; preparation of aryl and heteroaryl hydroxamic acids as inhibitors of histone deacetylase for treating proliferative diseases)

RN 603991-96-4 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-naphthalenylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)

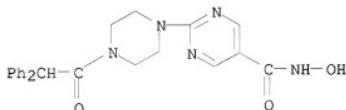
IT 603991-95-3P 603992-24-1P 603992-25-2P
603992-26-3P 603992-27-4P 603992-28-5P

RL: ARG (Analytical reagent use); PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate and reagent for detection/identification of histone deacetylase; preparation of aryl and heteroaryl hydroxamic acids as inhibitors of histone deacetylase for treating proliferative diseases)

RN 603991-95-3 CAPLUS

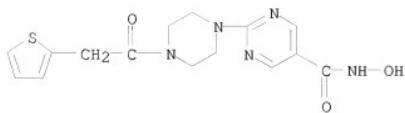
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RN 603992-24-1 CAPLUS

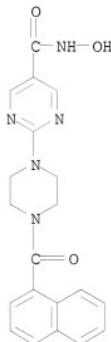
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-(2-thienyl)acetyl]-1-piperazinyl]- (CA INDEX NAME)

10/513699



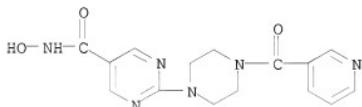
RN 603992-25-2 CAPLUS

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RN 603992-26-3 CAPLUS

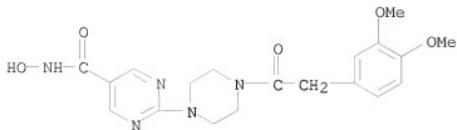
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(3-pyridinylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)



RN 603992-27-4 CAPLUS

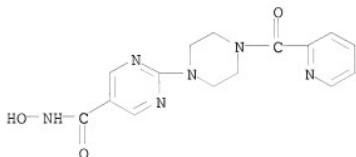
CN 5-Pyrimidinecarboxamide, 2-[4-[2-(3,4-dimethoxyphenyl)acetyl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

10/513699



RN 603992-28-5 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-pyridinylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)



REFERENCE COUNT:

6

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

10/513699

=> file erg
'ERG' IS NOT A VALID FILE NAME
SESSION CONTINUES IN FILE 'CAPLUS'
Enter "HELP FILE NAMES" at an arrow prompt (>) for a list of files
that are available. If you have requested multiple files, you can
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=> file reg		
COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	71.33	671.99
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	-10.40	-19.20

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STRUCTURE FILE UPDATES: 14 SEP 2008 HIGHEST RN 1049627-95-3
DICTIONARY FILE UPDATES: 14 SEP 2008 HIGHEST RN 1049627-95-3

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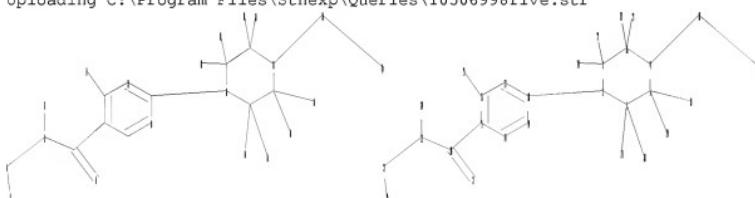
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<http://www.cas.org/support/stndoc/properties.html>

=>
Uploading C:\Program Files\Stnexp\Queries\10506998five.str



10/513699

chain nodes :
10 11 20 21 22 23 24 25 27 28 29 30 31 32 33 34 35
ring nodes :
1 2 3 4 5 14 15 16 17 18 19 26
chain bonds :
1-27 1-28 2-18 3-33 3-34 4-10 5-29 5-30 10-11 15-20 16-35 20-21 20-22
22-23 22-24 23-25 26-31 26-32
ring bonds :
1-2 1-5 2-3 3-26 4-5 4-26 14-15 14-19 15-16 16-17 17-18 18-19
exact/norm bonds :
1-2 1-5 2-3 2-18 3-26 4-10 4-5 4-26 10-11 20-21 20-22 22-23
exact bonds :
1-27 1-28 3-33 3-34 5-29 5-30 15-20 16-35 22-24 23-25 26-31 26-32
normalized bonds :
14-15 14-19 15-16 16-17 17-18 18-19
isolated ring systems :
containing 1 :

G1:C,N

G2:Ak,NH2,NO2

G3:O

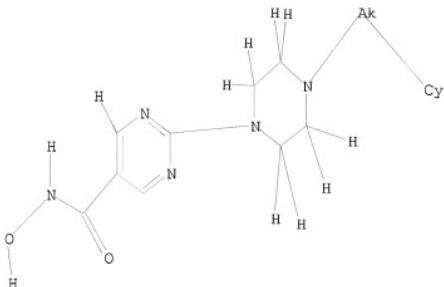
G4

G5:C,N,Zn,H

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 10:CLASS 11:Atom 14:Atom 15:Atom
16:Atom 17:Atom 18:Atom 19:Atom 20:CLASS 21:CLASS 22:CLASS 23:CLASS
24:CLASS 25:CLASS 26:Atom 27:CLASS 28:CLASS 29:CLASS 30:CLASS 31:CLASS
32:CLASS 33:CLASS 34:CLASS 35:CLASS

L12 STRUCTURE UPLOADED

=> d l12
L12 HAS NO ANSWERS
L12 STR



G1 C,N
 G2 Ak,NH₂,NO₂
 G3 O
 G4
 G5 C,N,Zn,H

Structure attributes must be viewed using STN Express query preparation.

=> s 112 full
 FULL SEARCH INITIATED 16:08:56 FILE 'REGISTRY'
 FULL SCREEN SEARCH COMPLETED - 1679 TO ITERATE

100.0% PROCESSED 1679 ITERATIONS 89 ANSWERS
 SEARCH TIME: 00.00.01

L13 89 SEA SSS FUL L12

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COST IN U.S. DOLLARS	ENTRY	SESSION
FULL ESTIMATED COST	178.82	850.81
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FILE 'CAPLUS' ENTERED AT 16:09:04 ON 15 SEP 2008
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FILE COVERS 1907 - 15 Sep 2008 VOL 149 ISS 12
FILE LAST UPDATED: 14 Sep 2008 (20080914/ED)

Caplus now includes complete International Patent Classification (IPC)
reclassification data for the second quarter of 2008.

Effective October 17, 2005, revised CAS Information Use Policies apply.
They are available for your review at:

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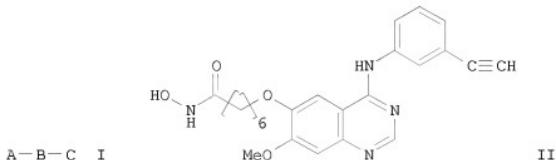
=> s l13 full
L14 9 L13

=> d ibib abs hitstr tot

L14 ANSWER 1 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2008:353001 CAPLUS
 DOCUMENT NUMBER: 148:355828
 TITLE: Multi-functional small molecules as anti-proliferative agents and their preparation
 INVENTOR(S): Cai, Xiong; Qian, Changgeng; Gould, Stephen; Zhai, Haixiao
 PATENT ASSIGNEE(S): Curis, Inc., USA
 SOURCE: PCT Int. Appl., 494pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008033747	A2	20080320	WO 2007-US77971	20070910
WO 2008033747	A9	20080724		
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RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA				
US 20080221132	A1	20080911	US 2007-852458	20070910
PRIORITY APPLN. INFO.:			US 2006-843590P	P 20060911
			US 2007-895889P	P 20070320

OTHER SOURCE(S): MARPAT 148:355828
 GI



AB The invention relates to the compns., methods, and applications of an approach to selective inhibition of several cellular or mol. targets with a single small mol. More specifically, the present invention relates to multi-functional small mols. of formula I wherein one functionality is capable of inhibiting histone deacetylases (HDAC) and the other functionality is capable of inhibiting a different cellular or mol. pathway involved in aberrant cell proliferation, differentiation or

survival. Compds. of formula I wherein A is a pharmacophore of an anticancer agent capable of inhibiting at least one cellular or mol. pathway involved in the aberrant cell proliferation, differentiation or survival; B is a linker; C is a zinc-binding moiety; and their geometrical isomers, enantiomers, diastereoisomers, racemates, pharmaceutically acceptable salts, prodrugs and solvates thereof, are claimed. Example compound II was prepared by a multistep procedure (procedure given). All the invention compds. were evaluated for their antiproliferative activity (some data given).

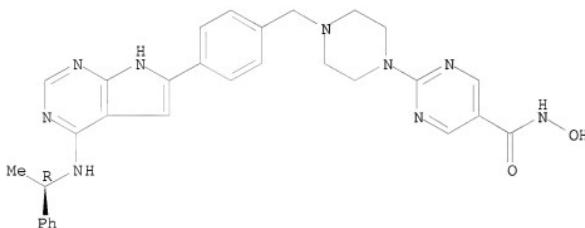
IT 1011716-90-7P

RL: PAC (Pharmacological activity); PKT (Pharmacokinetics); PRPH (Prophetic); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (prophetic starting material; preparation of multi-functional small mols. as antiproliferative agents)

RN 1011716-90-7 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[4-[4-[(1R)-1-phenylethyl]amino]-7H-pyrrolo[2,3-d]pyrimidin-6-yl]phenyl]methyl]-1-piperazinyl- (CA INDEX NAME)

Absolute stereochemistry.



L14 ANSWER 2 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2008:351928 CAPLUS
 DOCUMENT NUMBER: 148:355814
 TITLE: Preparation of (aralkylamino)(phenyl)pyrrolo[2,3-d]pyrimidine derivatives for use as protein tyrosine kinase (PTK) inhibitors
 INVENTOR(S): Cai, Xiong; Qian, Changgeng; Gould, Stephen
 PATENT ASSIGNEE(S): Curis, Inc., USA
 SOURCE: PCT Int. Appl., 123pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008033745	A2	20080320	WO 2007-US77968	20070910
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
US 20080161320	A1	20080703	US 2007-852440	20070910
PRIORITY APPLN. INFO.:			US 2006-843646P	P 20060911
			US 2007-895894P	P 20070320
OTHER SOURCE(S):	MARPAT	148:355814		
GI				

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB Fused bicyclic pyrimidine derivs. I and II [Ar = aryl, substituted arylheteroaryl or heteroaryl; Q = absent or (un)substituted alkyl; X = O, S, NH, or alkylamino; Z = O, S, NR1; Y = N or CR2; B = linker; D = C(O)NH2, NHC(S)CH3, CHC(O)NHacyl, etc.; R1 = H or (un)substituted alkyl; R2 = halo, (un)substituted aliphatic, aryl or heteroaryl], and their pharmaceutically acceptable salts, are prepared and disclosed as protein tyrosine kinase (PTK) inhibitors. Thus, e.g., III was prepared by N-alkylation of 1,4-dioxa-8-azaspiro[4.5]decane with 6-(4-(chloromethyl)phenyl)-N-((R)-1-phenylethyl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine (preparation given) and deprotection followed by condensation with 6-aminohexanoic acid Me ester and amidation with hydroxylamine. Select I were evaluated in EGFR assays, e.g., III demonstrated an IC50 value of ≤ 0.1 (μ M).

IT 1011716-90-7P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES

10/513699

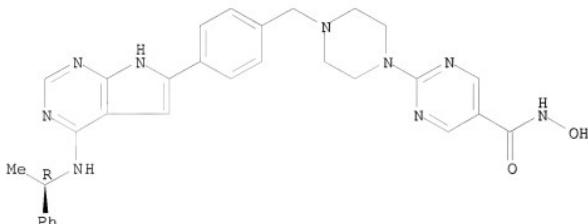
(Uses)

(preparation of (aralkylamino)(phenyl)pyrrolopyrimidine derivs. for use as protein tyrosine kinase (PTK) inhibitors)

RN 1011716-90-7 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[4-[(1R)-1-phenylethyl]amino]-7H-pyrrolo[2,3-d]pyrimidin-6-yl]phenyl]-1-piperazinyl- (CA INDEX NAME)

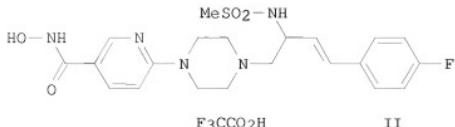
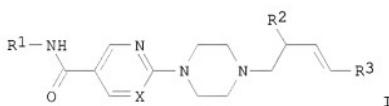
Absolute stereochemistry.



L14 ANSWER 3 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2007:816930 CAPLUS
 DOCUMENT NUMBER: 147:211903
 TITLE: Preparation of pyrimidine derivatives as histone deacetylase inhibitors
 INVENTOR(S): Marconnet-Decrane, Laurence Francoise Bernadette; Gaurrand, Sandrine Francoise Dominique; Angibaud, Patrick Rene
 PATENT ASSIGNEE(S): Janssen Pharmaceutica N.V., Belg.
 SOURCE: PCT Int. Appl., 62pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007082874	A1	20070726	WO 2007-EP50371	20070116
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JE, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, IJ, TM				
CA 2630717	A1	20070726	CA 2007-2630717	20070116
PRIORITY APPLN. INFO.:			EP 2006-100570	A 20060119
			WO 2007-EP50371	W 20070116

OTHER SOURCE(S): MARPAT 147:211903
 GI



AB The title compds. with general formula I [wherein R1 = OH or substituted phenyl; X = N or CH; R2 = amino, alkylamino, alkoxy, OH, etc.; R3 = (un)substituted Ph, naphthalene, or heterocycle] or N-oxide forms, pharmaceutically acceptable salts, or stereoisomeric forms thereof were prepared as histone deacetylase (HDAC) inhibitors for the treatment of proliferative diseases. For example, compound II was prepared in a multi-step synthesis. In vitro assay for inhibition of HDAC was performed to measure the inhibition of HDAC enzymic activity, and colorimetric assay was performed to determine cellular activity on A2780 tumor cells. II showed HDAC inhibitory and anti-proliferative activities in the above two assays with pIC50 values of 7.0 and 5.3, resp. Formulations containing I as active ingredients were also reported.

IT 944738-91-4P 944738-94-7P 944738-97-0P

944739-00-8P 944739-08-6P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of pyrimidine derivs. as histone deacetylase inhibitors)

RN 944738-91-4 CAPLUS

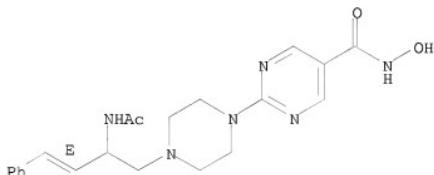
CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(acetylamino)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944738-90-3

CMF C21 H26 N6 O3

Double bond geometry as shown.



CM 2

CRN 76-05-1

CMF C2 H F3 O2



10/513699

RN 944738-94-7 CAPLUS

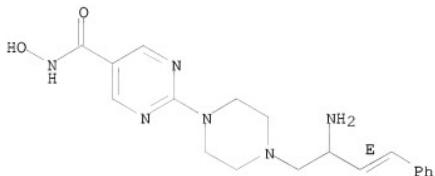
CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-amino-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944738-93-6

CMF C19 H24 N6 O2

Double bond geometry as shown.



CM 2

CRN 76-05-1

CMF C2 H F3 O2



RN 944738-97-0 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(2,5-dioxo-1-pyrrolidinyl)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

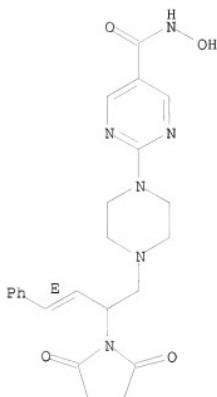
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CRN 944738-96-9

CMF C23 H26 N6 O4

Double bond geometry as shown.

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2



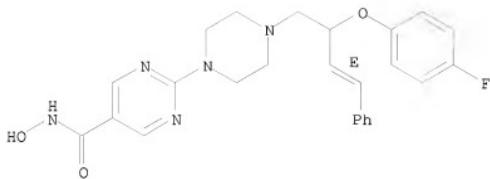
RN 944739-00-8 CAPLUS
CN 5-Pyrimidinecarboxamide, 2-[4-[(3E)-2-(4-fluorophenoxy)-4-phenyl-3-buten-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944738-99-2
CMF C25 H26 F N5 O3

Double bond geometry as shown.

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2

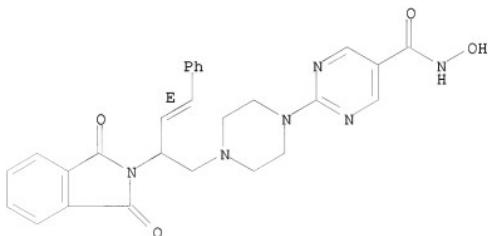


RN 944739-08-6 CAPLUS
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CRN 944739-07-5
CMF C27 H26 N6 O4

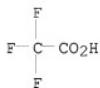
Double bond geometry as shown.



CM 2

10/513699

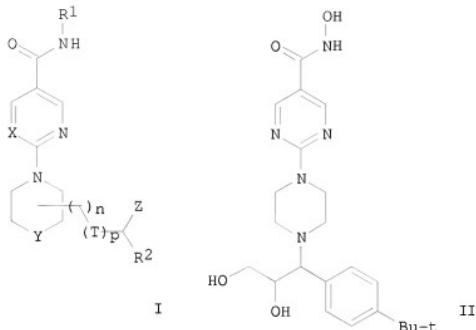
CRN 76-05-1
CMF C2 H F3 O2



REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 4 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2007:816806 CAPLUS
 DOCUMENT NUMBER: 147:211902
 TITLE: Preparation of pyrimidine derivatives as histone deacetylase inhibitors
 INVENTOR(S): Angibaud, Patrick Rene; Van Brandt, Sven Franciscus Anna; Marconnet-Decrane, Laurence Francoise Bernadette; Roux, Bruno
 PATENT ASSIGNEE(S): Janssen Pharmaceutica N.V., Belg.
 SOURCE: PCT Int. Appl., 63pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007082880	A1	20070726	WO 2007-EP50379	20070116
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JE, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
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PRIORITY APPLN. INFO.: EP 2006-100571				A 20060119
OTHER SOURCE(S): MARPAT 147:211902				
GI				



AB The title compds. with general formula I (wherein R1 = OH or substituted phenyl; R2 = -CH2OH, -CH2COH3, -CH2COCH2CH3, or -CH2CH(OH)CH2OH; T = N(R3), where R3 = H, alkyl, cycloalkyl, etc.; X = N or CH; Y = O, NH, CH2, etc.; n = 0-1; p = 0-1, provided that when p = 0 then n = 0 and Y = N, and -CH(R2)-Z is attached to Y; Z = (un)substituted aryl or heteroaryl) or N-oxide forms, pharmaceutically acceptable salts, or stereoisomeric forms thereof were prepared as histone deacetylase (HDAC) inhibitors for the treatment of proliferative diseases. For example, compound II was prepared in a multi-step synthesis. In vitro assay for inhibition of HDAC was performed to measure the inhibition of HDAC enzymic activity, and colorimetric assay was performed to determine cellular activity on A2780 tumor cells. II showed HDAC inhibitory and anti-proliferative activities in the above two assays with pIC50 values of 7.0 and 7.1, resp. Formulations containing I as active ingredients were also reported.

IT 944712-03-2P 944712-05-4P 944712-07-6P

944712-09-8P 944712-10-1P 944712-12-3P

$944712-14=5P$ $944712-16=7P$ $944712-18=9P$

BT: PAC (Pharmacological activity): SPM

RE: PAC (Pharmacological activity); SIR (Synthetic preparation); THER (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of pyrimidine derivs. as histone deacetylase inhibitors)

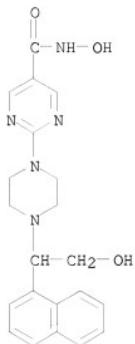
RN 944712-03-2 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-hydroxy-1-(1-naphthalenyl)ethyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-02-1
CMF C21 H23 N5 Q3

10/513699



CM 2

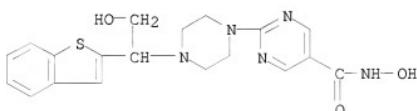
CRN 76-05-1
CMF C₂ H F₃ O₂



RN 944712-05-4 CAPLUS
CN 5-Pyrimidinecarboxamide, 2-[4-(1-benzo[b]thien-2-yl-2-hydroxyethyl)-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-04-3
CMF C₁₉ H₂₁ N₅ O₃ S



CM 2

10/513699

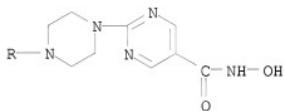
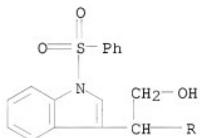
CRN 76-05-1
CMF C2 H F3 O2



RN 944712-07-6 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-hydroxy-1-[1-(phenylsulfonyl)-1H-indol-3-yl]ethyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-06-5
CMF C25 H26 N6 O5 S



CM 2

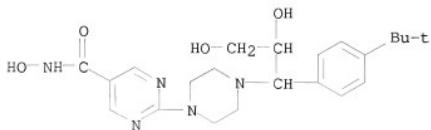
CRN 76-05-1
CMF C2 H F3 O2



RN 944712-09-8 CAPLUS
CN 5-Pyrimidinecarboxamide, 2-[4-[1-[4-(1,1-dimethylethyl)phenyl]-2,3-

10/513699

dihydroxypropyl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



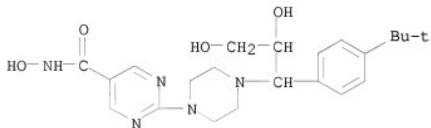
RN 944712-10-1 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1-[4-(1,1-dimethylethyl)phenyl]-2,3-dihydroxypropyl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1)
(CA INDEX NAME)

CM 1

CRN 944712-09-8

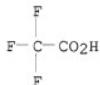
CMF C22 H31 N5 O4



CM 2

CRN 76-05-1

CMF C2 H F3 O2



RN 944712-12-3 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1(R,2S)-1-[4-(1,1-dimethylethyl)phenyl]-2,3-dihydroxypropyl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?)
(CA INDEX NAME)

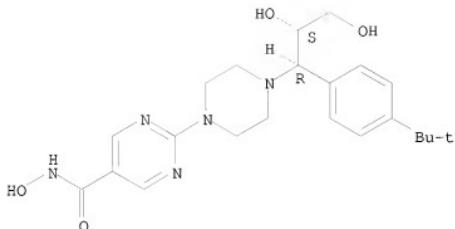
CM 1

CRN 944712-11-2

CMF C22 H31 N5 O4

Absolute stereochemistry.

10/513699



CM 2

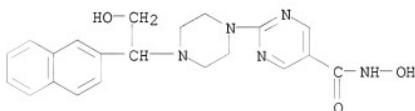
CRN 76-05-1
CMF C2 H F3 O2



RN 944712-14-5 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-hydroxy-1-(2-naphthalenyl)ethyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 944712-13-4
CMF C21 H23 N5 O3



CM 2

CRN 76-05-1
CMF C2 H F3 O2

10/513699

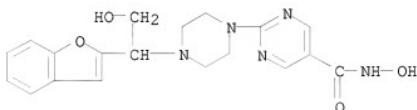


RN 944712-16-7 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1-(2-benzofuranyl)-2-hydroxyethyl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-15-6
CMF C19 H21 N5 O4



CM 2

CRN 76-05-1
CMF C2 H F3 O2



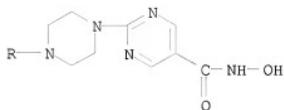
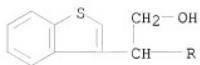
RN 944712-18-9 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-(1-benzo[b]thien-3-yl-2-hydroxyethyl)-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:?) (CA INDEX NAME)

CM 1

CRN 944712-17-8
CMF C19 H21 N5 O3 S

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2

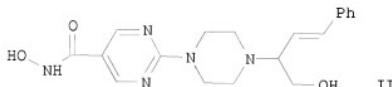
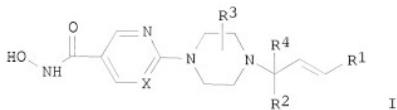


REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 5 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2006:101446 CAPLUS
 DOCUMENT NUMBER: 144:192266
 TITLE: Preparation of substituted propenyl piperazine derivatives as novel inhibitors of histone deacetylase
 INVENTOR(S): Van Brandt, Sven Franciscus Anna; Van Emelen, Kristof;
 Angibaud, Patrick Rene; Marconnet-Decrane, Laurence
 Francoise Bernadette; Arts, Janine
 PATENT ASSIGNEE(S): Janssen Pharmaceutica N.V., Belg.
 SOURCE: PCT Int. Appl., 67 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006010749	A2	20060202	WO 2005-EP53611	20050725
WO 2006010749	A3	20060608		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
AU 2005266311	A1	20060202	AU 2005-266311	20050725
CA 2572971	A1	20060202	CA 2005-2572971	20050725
EP 1776358	A2	20070425	EP 2005-777776	20050725
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, YU				
CN 1993356	A	20070704	CN 2005-80025487	20050725
JP 2008508234	T	20080321	JP 2007-523072	20050725
BR 2005013891	A	20080520	BR 2005-13891	20050725
KR 2007043978	A	20070426	KR 2007-701641	20070123
US 20070135424	A1	20070614	US 2007-626215	20070123
IN 2007DN00658	A	20070803	IN 2007-DN658	20070124
MX 200701119	A	20070315	MX 2007-1119	20070126
NO 200701117	A	20070227	NO 2007-1117	20070227
PRIORITY APPLN. INFO.:			EP 2004-77171	A 20040728
			US 2004-592357P	P 20040729
			WO 2005-EP53611	W 20050725

OTHER SOURCE(S): CASREACT 144:192266; MARPAT 144:192266
 GI



AB Substituted propenyl piperazine derivs. I, wherein X is independently N or CH; R1 is Ph, naphthalenyl or heterocyclyl; wherein each of said Ph or naphthalenyl is optionally substituted with one or two substituents each independently selected from halo, alkyl, alkyloxy, poly-halo-alkyl, aryl, hydroxy, cyano, amino, alkylcarbonylamino, alkylsulfonylamino, hydroxycarbonyl, alkyloxycarbonyl, hydroxyalkyl, alkyloxymethyl, aminomethyl, alkylaminomethyl, alkylcarbonylaminomethyl, alkylsulfonylaminomethyl, aminosulfonyl, alkylaminosulfonyl or heterocyclyl; R2 is hydrogen, -CH₂R₅, trifluoromethyl, -C(O)-R₆, or -CH-NR/R₈; wherein each R₅ is independently hydrogen, hydroxy, alkyloxy, alkyloxyalkyloxy, alkylcarbonyloxy, piperazinyl, N-methylpiperazinyl, morpholinyl, thiomorpholinyl, imidazolyl or triazolyl; each R₆ is independently hydroxy, alkyloxy, amino or mono- or di(alkyl)amino, cycloalkylamino, hydroxylalkylamino, piperazinyl, N-methylpiperazinyl, morpholinyl or thiomorpholinyl; each R₇ and R₈ are independently hydrogen, alkyl, alkylcarbonyl, alkylsulfonyl, or mono- or di(alkyl)aminosulfonyl; R₃ is hydrogen, hydroxymethyl, aminomethyl or mono- or di(alkyl)aminomethyl; R₄ is hydrogen or alkyl; were prepared and having histone deacetylase inhibiting enzymic activity and to inhibit proliferative conditions, such as cancer and psoriasis. Thus, propenyl piperazine derivative II was prepared and tested in vitro and in nude mice as inhibitor of histone deacetylase and was better than R306465 after oral administration. P21 enzyme linked immunosorbent assay has been applied to determine the p21 protein expression level in human A2780 ovarian carcinoma cells. In vitro assay for inhibition of histone deacetylase is reported. P21 induction was measured as the consequence of DNA damage or as the consequence of histone deacetylase inhibition. Antiproliferative activity of title compds. was determined on A2780 cells (neg. log value of the IC₅₀, pIC₅₀ = 7.9-8.2).

IT

875138-85-5P 875138-87-7P 875138-88-8P
 875138-89-9P 875138-90-2P 875138-91-3P
 875138-93-5P 875138-94-6P 875138-98-0P
 875139-00-7P 875139-02-9P 875139-04-1P
 875139-06-3P 875139-07-4P 875139-09-6P
 875139-11-0P 875139-13-2P 875139-14-3P
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10/513699

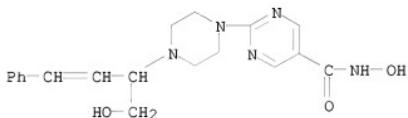
875139-70-1P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of substituted propenyl piperazine derivs. as novel inhibitors of histone deacetylase)

RN 875138-85-5 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(1-(hydroxymethyl)-3-phenyl-2-propen-1-yl)-1-piperazinyl]- (CA INDEX NAME)



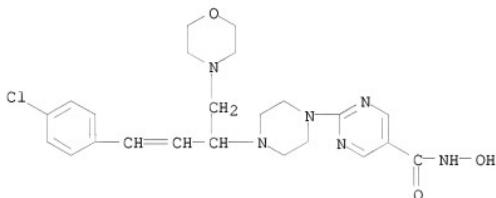
RN 875138-87-7 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3-(4-chlorophenyl)-1-(4-morpholinylmethyl)-2-propen-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-86-6

CMF C23 H29 Cl N6 O3



CM 2

CRN 76-05-1

CMF C2 H F3 O2

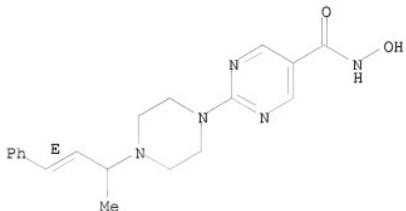


10/513699

RN 875138-88-8 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-methyl-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)

Double bond geometry as shown.



RN 875138-89-9 CAPLUS

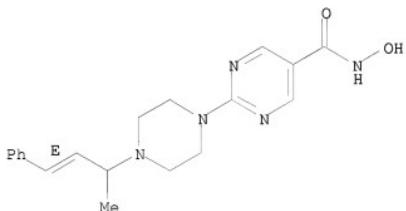
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-methyl-3-phenyl-2-propen-1-yl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-88-8

CMF C19 H23 N5 O2

Double bond geometry as shown.



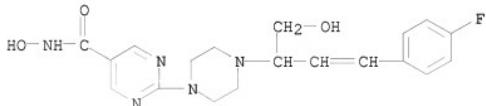
CM 2

CRN 76-05-1

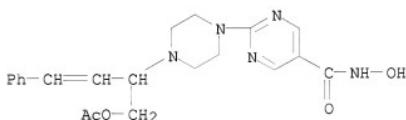
CMF C2 H F3 O2



RN 875138-90-2 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[3-(4-fluorophenyl)-1-(hydroxymethyl)-2-propen-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

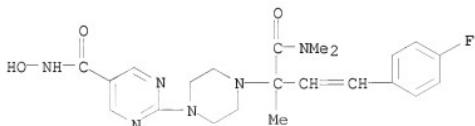


RN 875138-91-3 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(acetoxy)methyl]-3-phenyl-2-propen-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



RN 875138-93-5 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(dimethylamino)carbonyl]-3-(4-fluorophenyl)-1-methyl-2-propen-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-92-4
CMF C22 H27 F N6 O3

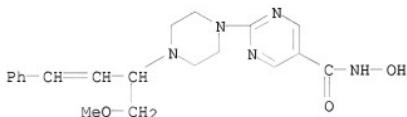
CM 2

10/513699

CRN 76-05-1
CMF C2 H F3 O2



RN 875138-94-6 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-(methoxymethyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)

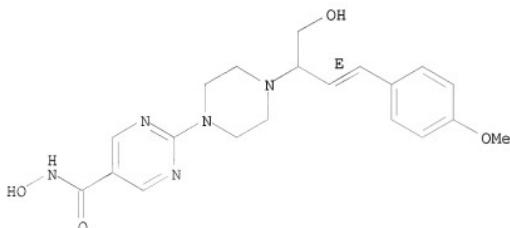


RN 875138-98-0 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-(hydroxymethyl)-3-(4-methoxyphenyl)-2-propen-1-yl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-97-9
CMF C20 H25 N5 O4

Double bond geometry as shown.



CM 2

CRN 76-05-1

10/513699

CMF C2 H F3 O2



RN 875139-00-7 CAPLUS

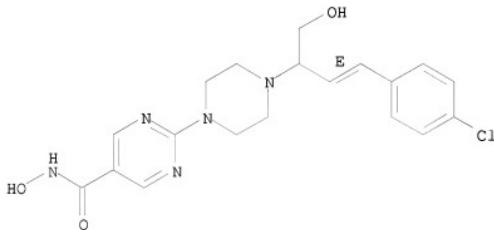
CN 5-Pyrimidinecarboxamide, 2-[4-[(2E)-3-(4-chlorophenyl)-1-(hydroxymethyl)-2-propen-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875138-99-1

CMF C19 H22 Cl N5 O3

Double bond geometry as shown.



CM 2

CRN 76-05-1

CMF C2 H F3 O2



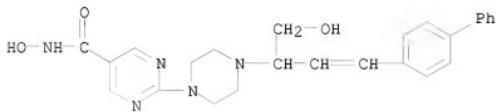
RN 875139-02-9 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[3-[1,1'-biphenyl]-4-yl]-1-(hydroxymethyl)-2-propen-1-yl]-1-piperazinyl-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

10/513699

CRN 875139-01-8
CMF C25 H27 N5 O3



CM 2

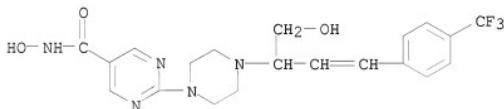
CRN 76-05-1
CMF C2 H F3 O2



RN 875139-04-1 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-(hydroxymethyl)-3-[4-(trifluoromethyl)phenyl]-2-propenyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-03-0
CMF C20 H22 F3 N5 O3



CM 2

CRN 76-05-1
CMF C2 H F3 O2

10/513699



RN 875139-06-3 CAPLUS

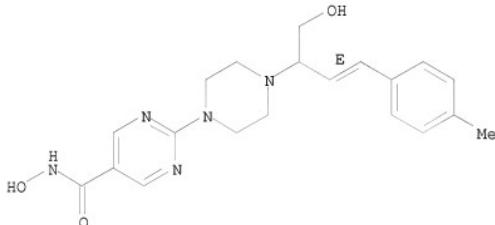
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(2E)-1-(hydroxymethyl)-3-(4-methylphenyl)-2-propen-1-yl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-05-2

CMF C20 H25 N5 O3

Double bond geometry as shown.



CM 2

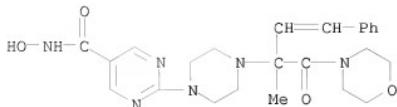
CRN 76-05-1

CMF C2 H F3 O2



RN 875139-07-4 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-methyl-1-(4-morpholinylcarbonyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)



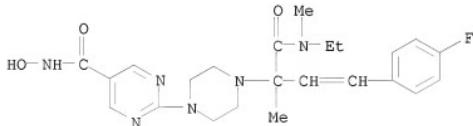
RN 875139-09-6 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(ethylmethylamino)carbonyl]-3-(4-fluorophenyl)-1-methyl-2-propenyl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-08-5

CMF C23 H29 F N6 O3



CM 2

CRN 76-05-1

CMF C2 H F3 O2



RN 875139-11-0 CAPLUS

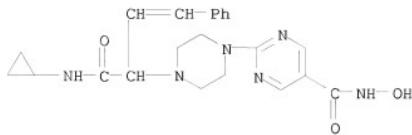
CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(cyclopropylamino)carbonyl]-3-phenyl-2-propenyl-1-yl]-1-piperazinyl]-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-10-9

CMF C22 H26 N6 O3

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2

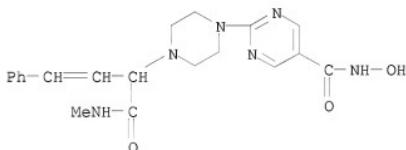


RN 875139-13-2 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(1-[(methylamino)carbonyl]-3-phenyl-2-propen-1-yl)-1-piperazinyl]-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-12-1
CMF C20 H24 N6 O3



CM 2

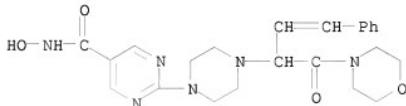
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10/513699



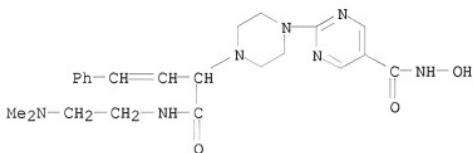
RN 875139-14-3 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-(4-morpholinylcarbonyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)



RN 875139-15-4 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[1-[(2-(dimethylamino)ethyl]amino]carbonyl]-3-phenyl-2-propen-1-yl]-1-piperazinyl-N-hydroxy- (CA INDEX NAME)



RN 875139-17-6 CAPLUS

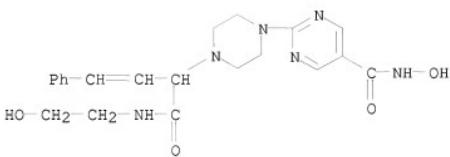
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[1-[(2-hydroxyethyl)amino]carbonyl]-3-phenyl-2-propen-1-yl]-1-piperazinyl-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-16-5

CMF C21 H26 N6 O4

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2

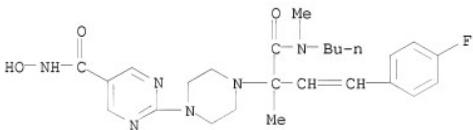


RN 875139-19-8 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-[(butylmethylamino)carbonyl]-3-(4-fluorophenyl)-2-propenyl-1-yl]-1-piperazinyl-N-hydroxy-, 2,2,2-trifluoroacetate (1:1) (CA INDEX NAME)

CM 1

CRN 875139-18-7
CMF C25 H33 F N6 O3

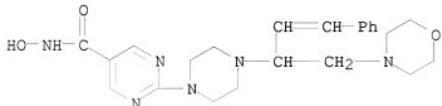


CM 2

CRN 76-05-1
CMF C2 H F3 O2

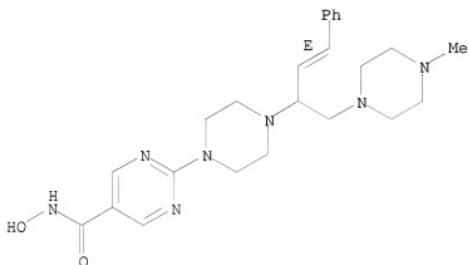


RN 875139-20-1 CAPLUS
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RN 875139-21-2 CAPLUS
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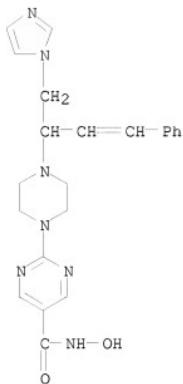
Double bond geometry as shown.



RN 875139-23-4 CAPLUS
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CM 1

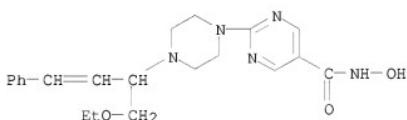
CRN 875139-22-3
 CMF C22 H25 N7 O2



CM 2

CRN 76-05-1
CMF C2 H F3 O2

RN 875139-24-5 CAPLUS
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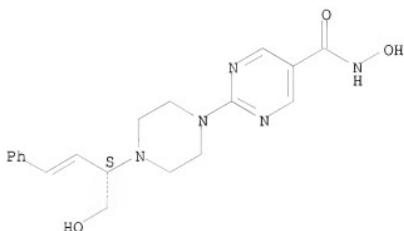


RN 875139-25-6 CAPLUS
 CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(1S)-1-(hydroxymethyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)

Absolute stereochemistry.

10/513699

Double bond geometry unknown.

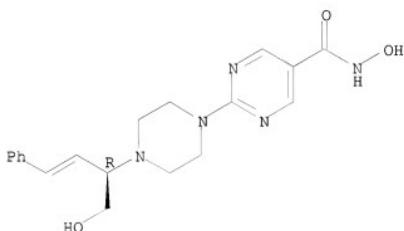


RN 875139-26-7 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(1R)-1-(hydroxymethyl)-3-phenyl-2-propen-1-yl]-1-piperazinyl]- (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry unknown.

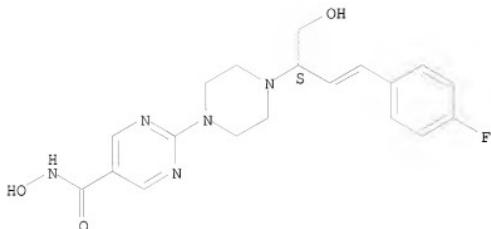


RN 875139-27-8 CAPLUS

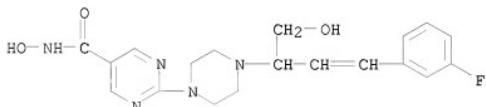
CN 5-Pyrimidinecarboxamide, 2-[4-[(1S)-3-(4-fluorophenyl)-1-(hydroxymethyl)-2-propen-1-yl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

Absolute stereochemistry.

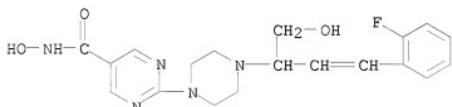
Double bond geometry unknown.



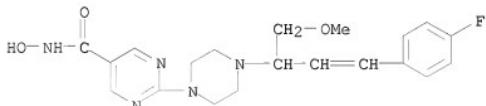
RN 875139-28-9 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[3-(3-fluorophenyl)-1-(hydroxymethyl)-2-propenyl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



RN 875139-29-0 CAPLUS
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RN 875139-30-3 CAPLUS
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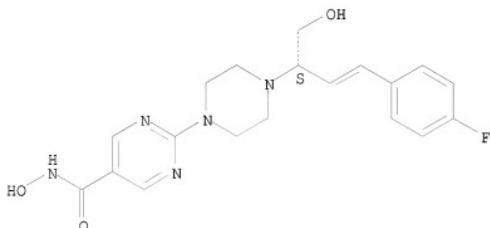


RN 875139-31-4 CAPLUS
 CN 5-Pyrimidinecarboxamide, 2-[4-[(1S)-3-(4-fluorophenyl)-1-(hydroxymethyl)-2-

10/513699

propen-1-yl]-1-piperazinyl]-N-hydroxy-, hydrochloride (1:1) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.

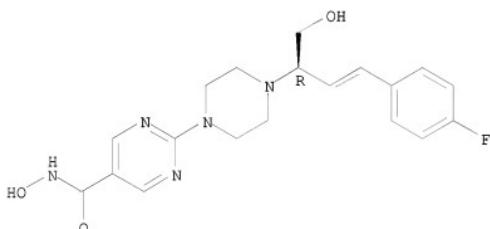


● HCl

RN 875139-69-8 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-((1R)-3-(4-fluorophenyl)-1-(hydroxymethyl)-2-propen-1-yl)-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.

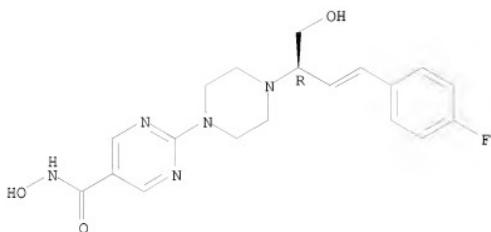


RN 875139-70-1 CAPLUS

CN 5-Pyrimidinecarboxamide, 2-[4-((1R)-3-(4-fluorophenyl)-1-(hydroxymethyl)-2-propen-1-yl)-1-piperazinyl]-N-hydroxy-, hydrochloride (1:1) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.

10/513699

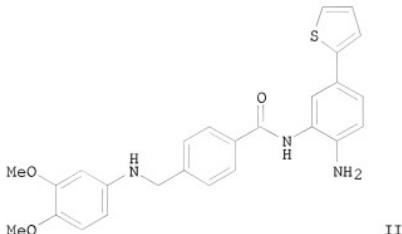
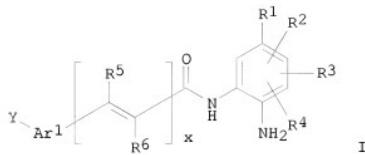


● HCl

L14 ANSWER 6 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2005:300395 CAPLUS
 DOCUMENT NUMBER: 142:355054
 TITLE: Preparation of amide derivatives as inhibitors of histone deacetylase
 INVENTOR(S): Moradei, Oscar; Paquin, Isabelle; Leit, Silvana;
 Frechette, Sylvie; Vaisburg, Arkadii; Besterman,
 Jeffrey M.; Tessier, Pierre; Mallais, Tammy C.
 PATENT ASSIGNEE(S): Methylgene, Inc., Can.
 SOURCE: PCT Int. Appl., 559 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005030705	A1	20050407	WO 2004-US31591	20040924
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RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2004276337	A1	20050407	AU 2004-276337	20040924
CA 2539117	A1	20050407	CA 2004-2539117	20040924
EP 1663953	A1	20060607	EP 2004-789074	20040924
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR				
CN 1882529	A	20061220	CN 2004-80034571	20040924
JP 2007506785	T	20070322	JP 2006-528279	20040924
US 20080132459	A1	20080605	US 2006-574088	20060323
JP 2008094847	A	20080424	JP 2007-281356	20071030
PRIORITY APPLN. INFO.:			US 2003-505884P	P 20030924
			US 2003-532973P	P 20031229
			US 2004-561082P	P 20040409
			JP 2006-528279	A3 20040924
			WO 2004-US31591	W 20040924

OTHER SOURCE(S): CASREACT 142:355054; MARPAT 142:355054
 GI



AB Title compds. I [Ar1 = (un)saturated-, (un)substituted-mono or fused poly-cyclic hydrocarbyl optionally containing 1-4 heteroatoms per ring; R1 = (un)substituted-mono-, -bi-, -tri-cyclic-aryl or -heteroaryl; R2, R3, and R4 independently = H, halo, amino, etc.; R5 and R6 independently = H, alkyl, aryl, etc.; x = 0-1; Y = any pharmaceutically acceptable chemical moiety consisting of 1 to 50 atoms with provisions] and their pharmaceutically acceptable salts, are prepared and disclosed as inhibitors of histone deacetylase. Thus, e.g., II was prepared by Suzuki coupling of 2-bromo-2-nitro-phenylamine (preparation given) with 2-thiopheneboronic acid followed by carbonylation with 4-[3,4-dimethoxy-(phenylamino)-methyl]benzoic acid (preparation given) and subsequent reduction. The inhibitory capability of I towards antiproliferative activity of histone deacetylase enzyme was evaluated using 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT) assay and it revealed that certain compds. of the invention had MTT IC 50 values in the range of below 1 up to 20 μ M. I as histone deacetylase inhibitors should prove useful in the treatment of diseases such as, but not limited to, cell proliferative disease, protozoal disease, and fungal disease.

IT 603985-86-0P 603985-88-2P 603985-90-6P
 603985-94-0P 603991-95-3P 603991-96-4P
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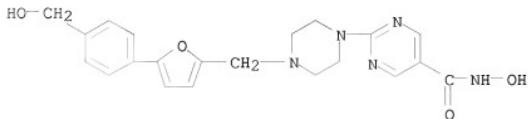
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of amide derivs. as inhibitors of histone deacetylase)

RN 603985-86-0 CAPLUS

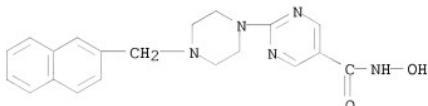
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CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(5-[4-(hydroxymethyl)phenyl]-2-furanyl)methyl]-1-piperazinyl]- (CA INDEX NAME)



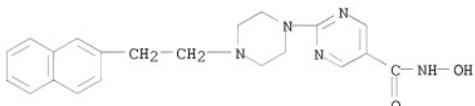
RN 603985-88-2 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-naphthalenylmethyl)-1-piperazinyl]- (CA INDEX NAME)



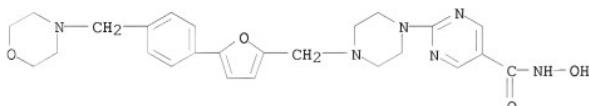
RN 603985-90-6 CAPLUS

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RN 603985-94-0 CAPLUS

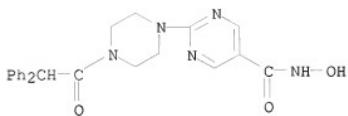
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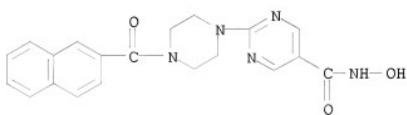
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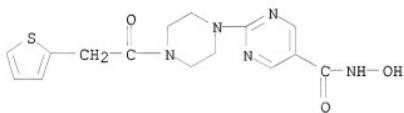
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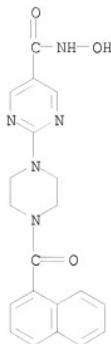
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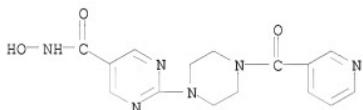
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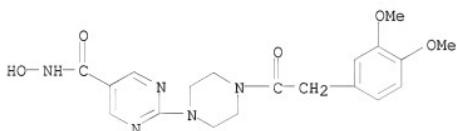
RN 603992-25-2 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(1-naphthalenylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)



RN 603992-26-3 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(3-pyridinylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)

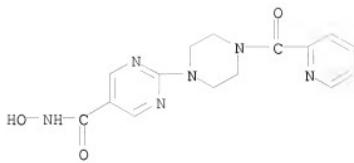


RN 603992-27-4 CAPLUS
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RN 603992-28-5 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-pyridinylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)

10/513699



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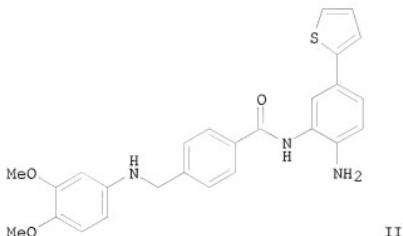
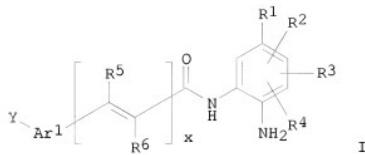
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THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 7 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2005:300394 CAPLUS
 DOCUMENT NUMBER: 142:373563
 TITLE: Preparation of amide derivatives as inhibitors of histone deacetylase
 INVENTOR(S): Moradei, Oscar; Paquin, Isabelle; Leit, Silvana;
 Frechette, Sylvie; Vaisburg, Arkadii; Besterman,
 Jeffrey M.; Tessier, Pierre; Mallais, Tammy C.
 PATENT ASSIGNEE(S): Methylgene, Inc., Can.
 SOURCE: PCT Int. Appl., 389 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005030704	A1	20050407	WO 2004-US31590	20040924
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, T2, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
JP 2008094847	A	20080424	JP 2007-281356 US 2003-505884P US 2003-532973P US 2004-561082P JP 2006-528279	20071030 P 20030924 P 20031229 P 20040409 A3 20040924
PRIORITY APPLN. INFO.:				

OTHER SOURCE(S): CASREACT 142:373563; MARPAT 142:373563
 GI



AB Title compds. I [Ar1 = (un)saturated-, (un)substituted-mono or fused poly-cyclic hydrocarbyl optionally containing 1-4 heteroatoms per ring; R1 = (un)substituted-mono-, -bi-, -tri-cyclic-aryl or -heteroaryl; R2, R3, and R4 independently = H, halo, amino, etc.; R5 and R6 independently = H, alkyl, aryl, etc.; x = 0-1; Y = any pharmaceutically acceptable chemical moiety consisting of 1 to 50 atoms with provisions] and their pharmaceutically acceptable salts, are prepared and disclosed as inhibitors of histone deacetylase. Thus, e.g., II was prepared by Suzuki coupling of 2-bromo-2-nitro-phenylamine (preparation given) with 2-thiopheneboronic acid followed by carbonylation with 4-[3,4-dimethoxy-(phenylamino)-methyl]benzoic acid (preparation given) and subsequent reduction. The inhibitory capability of I towards antiproliferative activity of histone deacetylase enzyme was evaluated using 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT) assay and it revealed that certain compds. of the invention had MTT IC₅₀ values in the range of below 1 up to 20 μM. I as histone deacetylase inhibitors should prove useful in the treatment of diseases such as, but not limited to, cell proliferative disease, protozoal disease, and fungal disease.

IT 603985-86-0P 603985-88-2P 603985-90-6P
 603985-94-0P 603991-95-3P 603991-96-4P
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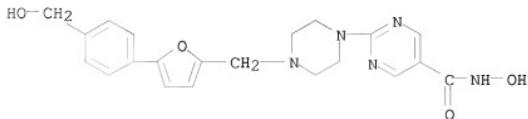
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of amide derivs. as inhibitors of histone deacetylase)

RN 603985-86-0 CAPLUS

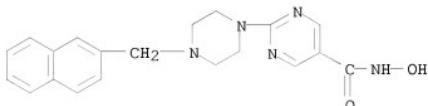
10/513699

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[(5-[4-(hydroxymethyl)phenyl]-2-furanyl)methyl]-1-piperazinyl]- (CA INDEX NAME)



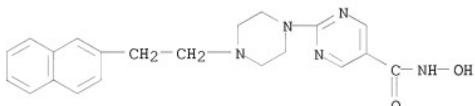
RN 603985-88-2 CAPLUS

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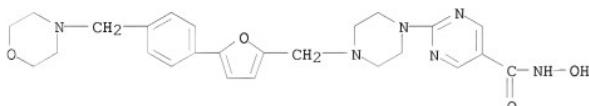
RN 603985-90-6 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-(2-naphthalenyl)ethyl]-1-piperazinyl]- (CA INDEX NAME)



RN 603985-94-0 CAPLUS

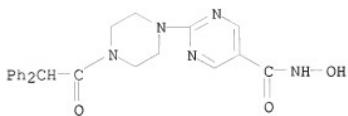
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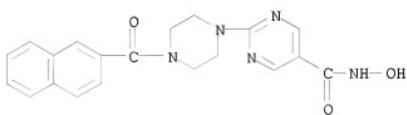
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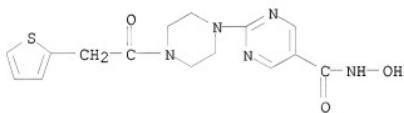
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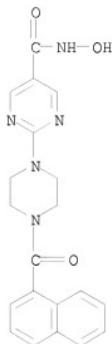
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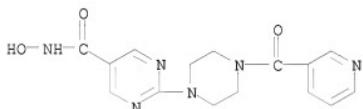
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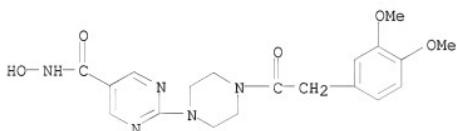
RN 603992-25-2 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(1-naphthalenylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)



RN 603992-26-3 CAPLUS
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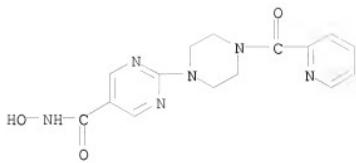


RN 603992-27-4 CAPLUS
CN 5-Pyrimidinecarboxamide, 2-[4-(2-(3,4-dimethoxyphenyl)acetyl)-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)



RN 603992-28-5 CAPLUS
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-pyridinylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)

10/513699



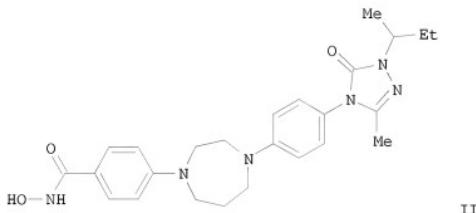
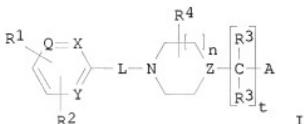
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THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 8 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2003:737723 CAPLUS
 DOCUMENT NUMBER: 139:261309
 TITLE: Preparation of N-hydroxy-5-piperazino(piperidino or diazepino)-2-pyrimidinecarboxamides and N-hydroxy-4-piperazino(piperidino or diazepino)benzamides as new inhibitors of histone deacetylase
 INVENTOR(S): Angibaud, Patrick Rene; Pilatte, Isabelle Noelle Constance; Van Brandt, Sven Franciscus Anna; Roux, Bruno; Ten Holte, Peter; Verdonck, Marc Gustaaf Celine; Meerpoel, Lieven; Dyatkin, Alexey Borisovich Janssen Pharmaceutica N.V., Belg.
 PATENT ASSIGNEE(S):
 SOURCE: PCT Int. Appl., 72 pp.
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 8
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003076400	A1	20030918	WO 2003-EP2514	20030311
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2475764	A1	20030918	CA 2003-2475764	20030311
AU 2003218736	A1	20030922	AU 2003-218736	20030311
EP 1485353	A1	20041215	EP 2003-711980	20030311
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
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CN 1639125	A	20050713	CN 2003-805675	20030311
CN 1642551	A	20050720	CN 2003-805833	20030311
NZ 534834	A	20050729	NZ 2003-534834	20030311
JP 2005526067	T	20050902	JP 2003-574621	20030311
CN 101007803	A	20070801	CN 2007-10005212	20030311
IN 2004DN02533	A	20070413	IN 2004-DN2533	20040831
US 20050107384	A1	20050519	US 2004-506998	20040908
ZA 2004007237	A	20050928	ZA 2004-7237	20040909
ZA 2004007235	A	20051004	ZA 2004-7235	20040909
ZA 2004007232	A	20051006	ZA 2004-7232	20040909
ZA 2004007233	A	20051006	ZA 2004-7233	20040909
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ZA 2004007236	A	20051006	ZA 2004-7236	20040909
MX 2004PA08806	A	20041126	MX 2004-PA8806	20040910
NO 2004004194	A	20041001	NO 2004-4194	20041001
PRIORITY APPLN. INFO.:				
		US 2002-363799P	P	20020313
		WO 2002-EP14833	A	20021223
		CN 2003-805921	A3	20030311



AB The title compds. [I; n = 0-3; t = 0-4; Q, X, Y = N, C; Z = N, CH; R1 = CONR7R8, NHCOR9, CO(alkanediyl)SR9, etc. (wherein R7, R8 = H, OH, alkyl, etc.; R9 = H, alkyl, alkylcarbonyl, etc.); R2 = H, halo, OH, etc.; L = a bond, alkanediyl, alkanediyoxy, NH, CO, NHCO; each R3 = H and one H atom can be replaced by aryl; R4 = H, OH, NH2, etc.; A = (un)substituted Ph, cyclohexyl, pyridyl, etc.], having histone deacetylase inhibiting enzymic activity, were prepared and formulated. E.g., a multi-step synthesis of II which showed pIC50 of 5.121 against HDAC, was given.

IT 603985-87-1P 603985-89-3P 603985-91-7P

603985-95-1P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of piperazino(piperidino or diazepino) substituted 2-pyrimidinecarbohydroxamic acids and N-hydroxybenzamides as new inhibitors of histone deacetylase)

RN 603985-87-1 CAPLUS

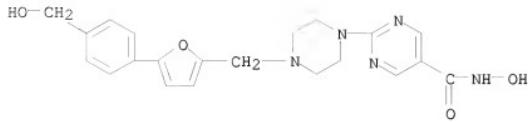
CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-([5-[4-(hydroxymethyl)phenyl]-2-furanyl)methyl]-1-piperazinyl-, 2,2,2-trifluoroacetate (5:4) (CA INDEX NAME)

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CRN 603985-86-0

CMF C21 H23 N5 O4

10/513699



CM 2

CRN 76-05-1
CMF C2 H F3 O2

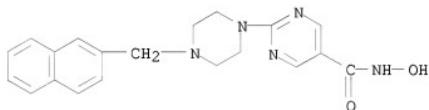


RN 603985-89-3 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-naphthalenylmethyl)-1-piperazinyl]-, 2,2,2-trifluoroacetate (5:4) (CA INDEX NAME)

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CRN 603985-88-2
CMF C20 H21 N5 O2



CM 2

CRN 76-05-1
CMF C2 H F3 O2



RN 603985-91-7 CAPLUS

<12/04/2007>

Erich Leese

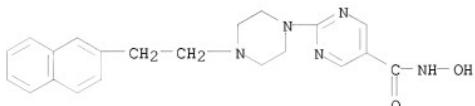
10/513699

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[2-(2-naphthalenyl)ethyl]-1-piperazinyl]-, 2,2,2-trifluoroacetate (5:4) (CA INDEX NAME)

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CRN 603985-90-6

CMF C21 H23 N5 O2



CM 2

CRN 76-05-1

CMF C2 H F3 O2



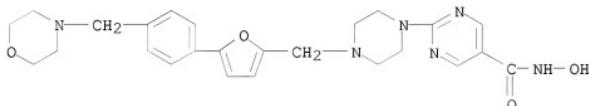
RN 603985-95-1 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-[5-[4-(4-morpholinylmethyl)phenyl]-2-furanyl]methyl]-1-piperazinyl-, 2,2,2-trifluoroacetate (1:2) (CA INDEX NAME)

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CRN 603985-94-0

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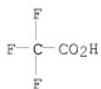


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10/513699



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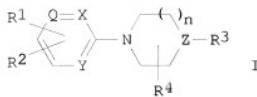
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THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 9 OF 9 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2003:737586 CAPLUS
 DOCUMENT NUMBER: 139:261308
 TITLE: Preparation of aryl and heteroaryl hydroxamic acids as inhibitors of histone deacetylase for treating proliferative diseases
 INVENTOR(S): Van Emelen, Kristof; Verdonck, Marc Gustaaf Celine;
 Van Brandt, Sven Franciscus Anna; Angibaud, Patrick
 Rene; Meerpoel, Lieven; Dyatkin, Alexey Borisovich
 Janssen Pharmaceutica N.V., Belg.
 PATENT ASSIGNEE(S):
 SOURCE: PCT Int. Appl., 52 pp.
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 8
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003075929	A1	20030918	WO 2003-EP2515	20030311
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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
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EP 1485099	A1	20041215	EP 2003-711981	20030311
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NZ 534832	A	20050930	NZ 2003-534832	20030311
CN 101007803	A	20070801	CN 2007-10005212	20030311
IN 2004DN02537	A	20070112	IN 2004-DN2537	20040831
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NO 2004004113	A	20040928	NO 2004-4113	20040928
PRIORITY APPLN. INFO.:				
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		WO 2002-EP14833	A	20021223
		CN 2003-805921	A3	20030311
		WO 2003-EP2515	W	20030311

OTHER SOURCE(S): MARPAT 139:261308



AB This invention comprises aryl and heteroaryl hydroxamic acids (shown as I; variables defined below; e.g. II) having histone deacetylase inhibiting enzymic activity; their preparation, compns. containing them and their use as a medicine. Compds. I show excellent in-vitro histone deacetylase inhibiting enzymic activity, have advantageous properties with regard to cellular activity and specific properties with regard to inhibition of cell cycle progression at both G1 and G2 checkpoints (p21 induction capacity), and show good metabolic stability and high bioavailability and more particular show oral bioavailability. They can also be used for detection and identification of histone deacetylase. General synthetic procedures and characterization data for twenty-seven I are included; also, preps. of 12 intermediates are included. For example, a 59 % yield of 2-(4-(dimethylaminosulfonyl)piperazin-1-yl)pyrimidine-5-carbohydroxamic acid was obtained by removing the O-tetrahydropyranyl group of its ester using trifluoroacetic acid; the ester was prepared in 61 % yield from N¹-(ethylcarboximidoyl)-N,N-dimethyl-3-propanediamine monohydrochloride, sodium 2-[4-(dimethylaminosulfonyl)piperazin-1-yl]pyrimidine-5-carboxylate, O-(tetrahydro-2H-pyran-2-yl)hydroxylamine, and 1-hydroxy-1H-benzotriazole in CH₂C₁₂/THF. The sodium salt was obtained by base hydrolysis of the Et ester; the ester was prepared in 73 % yield from Et 2-(piperazin-1-yl)pyrimidine-5-carboxylate and dimethylsulfamoyl chloride; Et 2-(piperazin-1-yl)pyrimidine-5-carboxylate was obtained in <96 % yield from Et 2-(4-benzylpiperazin-1-yl)pyrimidine-5-carboxylate by hydrogenation using Pd/C; the benzyl derivative was obtained from 1-(phenylmethyl)piperazine, (135 mL) was added gradually to a solution of potassium carbonate (0.18 mol) and 2-(methylsulfonyl)-5-pyrimidinecarboxylic acid Et ester, K₂CO₃ in MeCN. For I: n is 0-3; Q, X and Y are N or C; Z is N or CH; R1 is -C(O)NR5R6, -N(H)C(O)R7, -C(O)-C1-6alkanediylR7, -NR8C(O)N(OH)R7, -NR8C(O)C1-6alkanediylR7, -NR8C(O)C:N(OH)R7 or another Zn-chelating-group; R2 is H, halo, hydroxy, amino, nitro, Cl-6alkyl, Cl-6alkyloxy, trifluoromethyl, di(Cl-6-alkyl)amino, hydroxymino or naphthalenylsulfonylpiperazinyl. R3 is H, Cl-6-alkyl, arylC2-6alkanediyl, furanylcarbonyl, naphthalenylcarbonyl, -C(O)phenylR9, Cl-6alkylaminocarbonyl, aminosulfonyl, arylaminosulfonyl, aminosulfonylamino, di(Cl-6-alkyl)aminosulfonylamino, arylaminosulfonylamino, aminosulfonylaminoCl-6-alkyl, arylaminosulfonylaminoCl-6alkyl, di(Cl-6-alkyl)aminocl-6alkyl, Cl-12-alkylsulfonyl, di(Cl-6-alkyl)aminosulfonyl, trihaloCl-6-alkylsulfonyl, di(aryl)Cl-6alkylcarbonyl, thiophenylCl-6alkylcarbonyl, pyridinylcarbonyl or arylCl-6alkylcarbonyl. R4 is H, hydroxy, amino, hydroxyCl-6alkyl, Cl-6alkyl, Cl-6alkyloxy,

arylCl-6-alkyl, aminocarbonyl, hydroxycarbonyl, aminoCl-6-alkyl, aminocarbonylCl-6-alkyl, hydroxycarbonylCl-6-alkyl, hydroxyaminocarbonyl, Cl-6-alkyloxycarbonyl, Cl-6-alkylaminoCl-6-alkyl or di(Cl-6-alkyl)aminoCl-6-alkyl; when R3 and R4 are present on the same C atom, R3 and R4 together may form -C(O)-NH-CH₂-NR10- wherein R10 is H or aryl; when R3 and R4 are present on adjacent C atoms, R3 and R4 together may form :CH-CH:CH-CH: ; addnl. details are given in the claims.

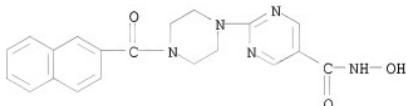
IT 603991-96-4P

RL: ARG (Analytical reagent use); PAC (Pharmacological activity); PKT (Pharmacokinetics); SPN (Synthetic preparation); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate and reagent for detection/identification of histone deacetylase; preparation of aryl and heteroaryl hydroxamic acids as inhibitors of histone deacetylase for treating proliferative diseases)

RN 603991-96-4 CAPLUS

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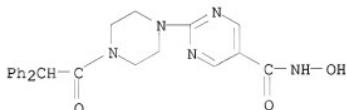
IT 603991-95-3P 603992-24-1P 603992-25-2P
603992-26-3P 603992-27-4P 603992-28-5P

RL: ARG (Analytical reagent use); PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate and reagent for detection/identification of histone deacetylase; preparation of aryl and heteroaryl hydroxamic acids as inhibitors of histone deacetylase for treating proliferative diseases)

RN 603991-95-3 CAPLUS

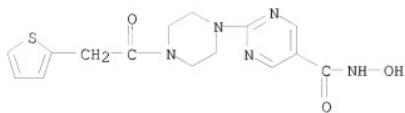
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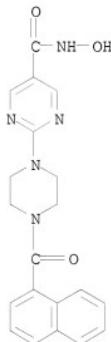
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10/513699



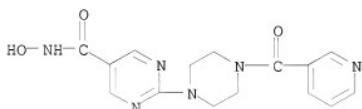
RN 603992-25-2 CAPLUS

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RN 603992-26-3 CAPLUS

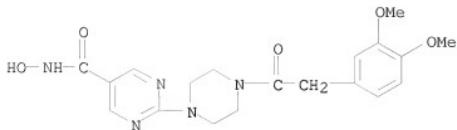
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RN 603992-27-4 CAPLUS

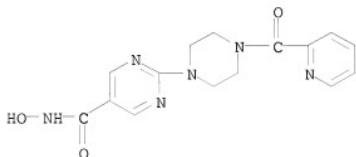
CN 5-Pyrimidinecarboxamide, 2-[4-[2-(3,4-dimethoxyphenyl)acetyl]-1-piperazinyl]-N-hydroxy- (CA INDEX NAME)

10/513699



RN 603992-28-5 CAPLUS

CN 5-Pyrimidinecarboxamide, N-hydroxy-2-[4-(2-pyridinylcarbonyl)-1-piperazinyl]- (CA INDEX NAME)



REFERENCE COUNT:

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THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

10/513699

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Connection closed by remote host